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(71) Applicant (for all designated States except US): **ASTRAZENECA AB** [SE/SE]; S-151 85 Södertälje (SE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **FINLAY, Raymond** [GB/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield, Cheshire SK10 4TG (GB). **TUCKER, Howard** [GB/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield, Cheshire SK10 4TG (GB). **WATERSON, David** [GB/GB]; AstraZeneca R & D Alderley, Alderley Park, Macclesfield, Cheshire SK10 4TG (GB).

(74) Agent: **GLOBAL INTELLECTUAL PROPERTY**; AstraZeneca AB, S-151 85 Södertälje (SE).

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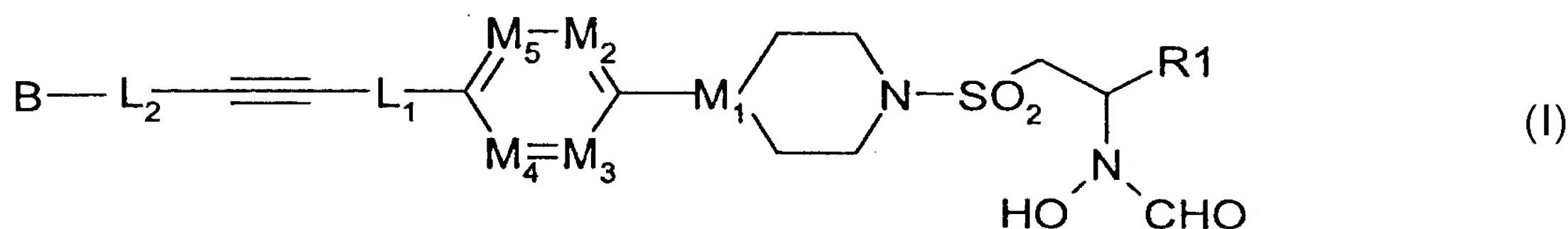
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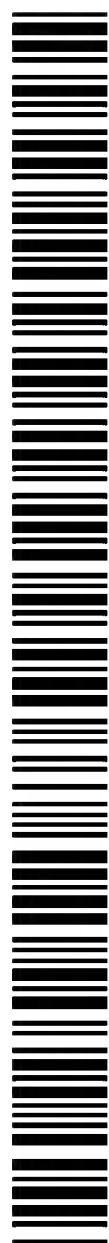
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(54) Title: ARYLPIPERAZINES AND ARYLPIPERIDINES AND THEIR USE AS METALLOPROTEINASE INHIBITING AGENTS



(57) Abstract: Compounds of the formula I useful as metalloproteinase inhibitors, especially as inhibitors of MMP 13.



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## ARYLPIPERAZINES AND ARYLPIPERIDINES AND THEIR USE AS METALLOPROTEINASE INHIBITING AGENTS

COMPOUNDS

The present invention relates to compounds useful in the inhibition of metalloproteinases and in particular to pharmaceutical compositions comprising these, as well as their use. In particular, the compounds of this invention are inhibitors of matrix metalloproteinase 13 (MMP13), known also as collagenase 3.

Metalloproteinases are a superfamily of proteinases (enzymes) whose numbers in recent years have increased dramatically. Based on structural and functional considerations these enzymes have been classified into families and subfamilies as described in N.M. Hooper (1994) FEBS Letters 354:1-6. Examples of metalloproteinases include the matrix metalloproteinases (MMPs); the reprotolysin or adamalysin or MDC family which includes the secretases and sheddases such as TNF converting enzymes (ADAM10 and TACE); the astacin family which include enzymes such as procollagen processing proteinase (PCP); and other metalloproteinases such as aggrecanase, the endothelin converting enzyme family and the angiotensin converting enzyme family.

Metalloproteinases are believed to be important in a plethora of physiological disease processes that involve tissue remodelling such as embryonic development, bone formation and uterine remodelling during menstruation. This is based on the ability of the metalloproteinases to cleave a broad range of matrix substrates such as collagen, proteoglycan and fibronectin. Metalloproteinases are also believed to be important in the processing, or secretion, of biological important cell mediators, such as tumour necrosis factor (TNF); and the post translational proteolysis processing, or shedding, of biologically important membrane proteins, such as the low affinity IgE receptor CD23 (for a more complete list see N. M. Hooper *et al.*, (1997) Biochem J. 321:265-279).

Metalloproteinases have been associated with many diseases or conditions. Inhibition of the activity of one or more metalloproteinases may well be of benefit in these diseases or conditions, for example: various inflammatory and allergic diseases such as, inflammation of the joint (especially rheumatoid arthritis, osteoarthritis and gout), inflammation of the gastro-intestinal tract (especially inflammatory bowel disease, ulcerative colitis and gastritis), inflammation of the skin (especially psoriasis, eczema, dermatitis); in tumour metastasis or invasion; in disease associated with uncontrolled

degradation of the extracellular matrix such as osteoarthritis; in bone resorptive disease (such as osteoporosis and Paget's disease); in diseases associated with aberrant angiogenesis; the enhanced collagen remodelling associated with diabetes, periodontal disease (such as gingivitis), corneal ulceration, ulceration of the skin, post-operative conditions (such as colonic anastomosis) and dermal wound healing; demyelinating diseases of the central and peripheral nervous systems (such as multiple sclerosis); Alzheimer's disease; extracellular matrix remodelling observed in cardiovascular diseases such as restenosis and atherosclerosis; and chronic obstructive pulmonary diseases, COPD (for example, the role of MMPs such as MMP12 is discussed in Anderson & Shinagawa, 1999, Current Opinion in Anti-inflammatory and Immunomodulatory Investigational Drugs, 1(1): 29-38).

The matrix metalloproteinases (MMPs) are a family of structurally-related zinc-containing endopeptidases which mediate the breakdown of connective tissue macromolecules. The mammalian MMP family is composed of at least twenty enzymes, classically divided into four sub-groups based on substrate specificity and domain structure [Alexander & Werb (1991) in Hay, E.D. ed. "Cell Biology of the Extracellular Matrix", New York, Plenum Press, 255-302; Murphy & Reynolds (1993) in Royce, P.M. & Steinman, B. eds. "Connective Tissue and its Heritable Disorders", New York, Wiley-Liss Inc., 287-316; Birkedal-Hansen (1995) Curr. Opin. Cell Biol. 7:728-735]. The sub-groups are the collagenases (such as MMP1, MMP8, MMP13), the stromelysins (such as MMP3, MMP10, MMP11), the gelatinases (such as MMP2, MMP9) and the membrane-type MMPs (such as MMP14, MMP15, MMP16, MMP17). Enzyme activity is normally regulated *in vivo* by tissue inhibitors of metalloproteinases (TIMPs).

Because of their central role in re-modelling connective tissue, both as part of normal physiological growth and repair and as part of disease processes, there has been substantial interest in these proteins as targets for therapeutic intervention in a wide range of degenerative and inflammatory diseases, such as arthritis, atherosclerosis, and cancer (Whittaker *et al* (1999) Chem. Rev. 99:2735-2776).

A number of MMP inhibitor compounds are known and some are being developed for pharmaceutical uses (see for example the review by Beckett & Whittaker (1998) Exp. Opin. Ther. Patents, 8(3):259-282). Different classes of compounds may have different

degrees of potency and selectivity for inhibiting various MMPs. Whittaker M. *et al* (1999, Chem. Rev. 99:2735-2776) review a wide range of known MMP inhibitor compounds. They state that an effective MMP inhibitor requires a zinc binding group or ZBG (functional group capable of chelating the active site zinc(II) ion), at least one functional group which provides a hydrogen bond interaction with the enzyme backbone, and one or more side chains which undergo effective van der Waals interactions with the enzyme subsites. Zinc binding groups in known MMP inhibitors include hydroxamic acids (-C(O)NHOH), reverse hydroxamates (-N(OH)CHO), thiols, carboxylates and phosphonic acids.

We have discovered a new class of compounds that are inhibitors of metalloproteinases and are of particular interest in inhibiting MMP13. The compounds of this invention have beneficial potency and/or pharmacokinetic properties. In particular they show selectivity for MMP13.

MMP13, or collagenase 3, was initially cloned from a cDNA library derived from a breast tumour [J. M. P. Freije *et al.* (1994) Journal of Biological Chemistry 269(24):16766-16773]. PCR-RNA analysis of RNAs from a wide range of tissues indicated that MMP13 expression was limited to breast carcinomas as it was not found in breast fibroadenomas, normal or resting mammary gland, placenta, liver, ovary, uterus, prostate or parotid gland or in breast cancer cell lines (T47-D, MCF-7 and ZR75-1). Subsequent to this observation MMP13 has been detected in transformed epidermal keratinocytes [N. Johansson *et al.*, (1997) Cell Growth Differ. 8(2):243-250], squamous cell carcinomas [N. Johansson *et al.*, (1997) Am. J. Pathol. 151(2):499-508] and epidermal tumours [K. Airola *et al.*, (1997) J. Invest. Dermatol. 109(2):225-231]. These results are suggestive that MMP13 is secreted by transformed epithelial cells and may be involved in the extracellular matrix degradation and cell-matrix interaction associated with metastasis especially as observed in invasive breast cancer lesions and in malignant epithelia growth in skin carcinogenesis.

Recent published data implies that MMP13 plays a role in the turnover of other connective tissues. For instance, consistent with MMP13's substrate specificity and preference for degrading type II collagen [P. G. Mitchell *et al.*, (1996) J. Clin. Invest. 97(3):761-768; V. Knauper *et al.*, (1996) The Biochemical Journal 271:1544-1550], MMP13 has been hypothesised to serve a role during primary ossification and skeletal



remodelling [M. Stahle-Backdahl *et al.*, (1997) Lab. Invest. 76(5):717-728; N. Johansson *et al.*, (1997) Dev. Dyn. 208(3):387-397], in destructive joint diseases such as rheumatoid and osteo-arthritis [D. Wernicke *et al.*, (1996) J. Rheumatol. 23:590-595; P. G. Mitchell *et al.*, (1996) J. Clin. Invest. 97(3):761-768; O. Lindy *et al.*, (1997) Arthritis Rheum 40(8):1391-1399]; and during the aseptic loosening of hip replacements [S. Imai *et al.*, (1998) J. Bone Joint Surg. Br. 80(4):701-710]. MMP13 has also been implicated in chronic adult periodontitis as it has been localised to the epithelium of chronically inflamed mucosa human gingival tissue [V. J. Uitto *et al.*, (1998) Am. J. Pathol 152(6):1489-1499] and in remodelling of the collagenous matrix in chronic wounds [M. Vaalamo *et al.*, (1997) J. Invest. Dermatol. 109(1):96-101].

US 6100266 and WO-99/38843 disclose compounds of the general formula

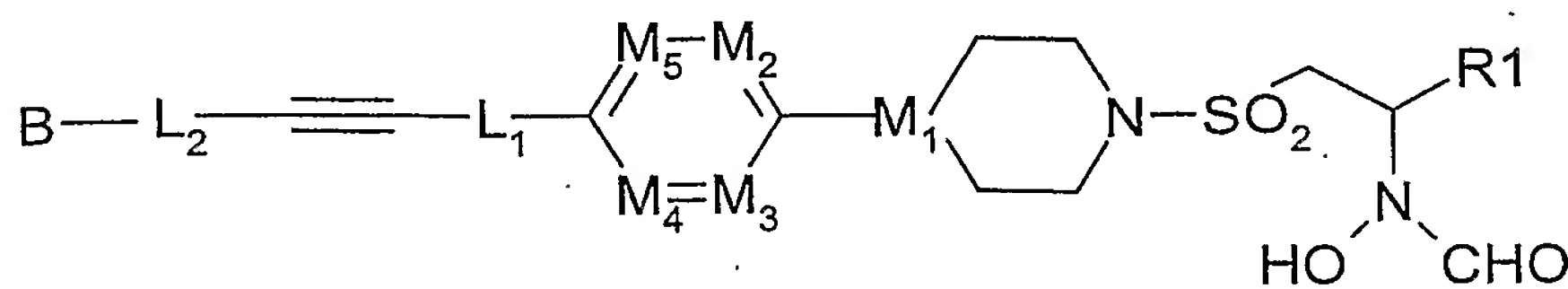


for use in the manufacture of a medicament for the treatment or prevention of a condition associated with matrix metalloproteinases. Specifically disclosed is the compound N-{1S-[4-(4-Chlorophenyl) piperazine-1-sulfonylmethyl]-2-methylpropyl}-N-hydroxyformamide.

WO-00/12478 discloses arylpiperazines that are matrix metalloproteinase inhibitors, including compounds with an hydroxamic acid zinc binding group and compounds with a reverse hydroxamate zinc binding group.

We have now discovered compounds that are potent MMP13 inhibitors and have desirable activity profiles.

In a first aspect of the invention we now provide a compound of the formula I



I

wherein

**B** is selected from H, C<sub>1-6</sub>alkyl, up to C12 cycloalkyl, up to C12 aryl, and up to C12 heteroaryl;

**B** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy;

**L**<sub>1</sub> and **L**<sub>2</sub> are each independently selected from a direct bond and C<sub>1-6</sub>alkyl;

**M**<sub>1</sub>, **M**<sub>2</sub>, **M**<sub>3</sub>, **M**<sub>4</sub> and **M**<sub>5</sub> are each independently selected from N and C;

**R**<sub>1</sub> is the group -**X**-**Y**;

**X** is C<sub>1-6</sub>alkyl;

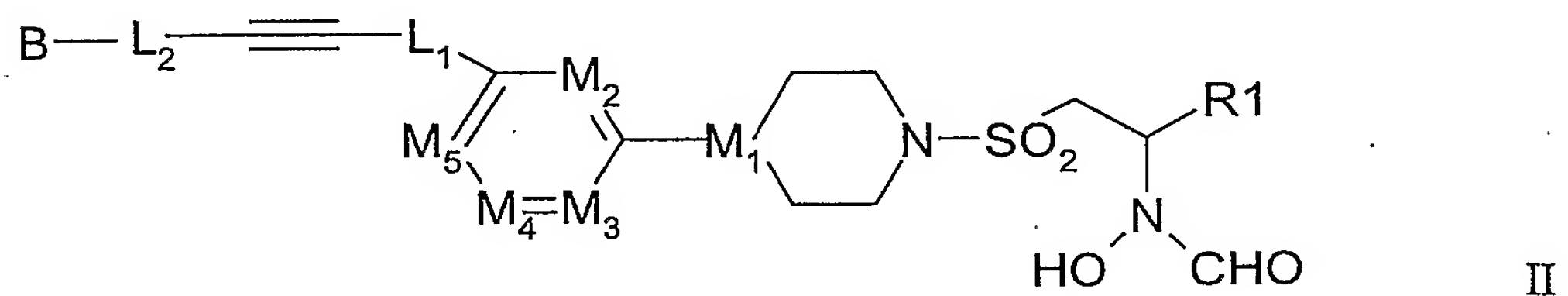
**Y** is selected from up to C10 cycloalkyl, up to C10 aryl, and up to C10 heteroaryl;

**Y** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy;

Any alkyl group outlined above may be straight chain or branched;

Any heteroaryl group outlined above is an aromatic ring containing one or more heteroatoms independently selected from N, O, S.

In a further aspect of the invention we provide compounds of the formula II



wherein **B**, **L**<sub>1</sub>, **L**<sub>2</sub>, **M**<sub>1</sub>, **M**<sub>2</sub>, **M**<sub>3</sub>, **M**<sub>4</sub>, **M**<sub>5</sub>, **R**<sub>1</sub>, **X** and **Y** are as defined above for the compound of formula I.

Preferred compounds of the formulae I or II are those wherein any one or more of the following apply:

**B** is selected from H, C<sub>1-6</sub>alkyl, C<sub>6</sub> aryl, and up to C<sub>6</sub> heteroaryl; preferably **B** is H, C<sub>2-4</sub>alkyl, C<sub>6</sub> aryl, and up to C<sub>6</sub> heteroaryl; most preferably **B** is up to C<sub>6</sub> heteroaryl;

**B** is not substituted or is substituted by at least one group selected from CF<sub>3</sub>, CN, halogen (preferably fluoro or chloro), C<sub>1-4</sub>alkyl;

At least one of  $L_1$  and  $L_2$  is a direct bond; preferably each of  $L_1$  and  $L_2$  is a direct bond;

$M_1$  is N;

At least one of  $M_2$ ,  $M_3$ ,  $M_4$ ,  $M_5$  is C, and at least one of  $M_2$ ,  $M_3$ ,  $M_4$ ,  $M_5$  is N;  
preferably each of  $M_4$  and  $M_5$  is C, and at least one of  $M_2$  and  $M_3$  is N; most preferably each of  $M_4$  and  $M_5$  is C,  $M_2$  is C or N and  $M_3$  is N;

X is  $C_{2-5}$ alkyl; preferably X is  $C_{2-3}$ alkyl;

Y is  $C_6$  aryl or  $C_6$  heteroaryl; preferably Y is phenyl, pyridyl, pyrimidinyl, or pyrazinyl; most preferably Y is pyrimidinyl;

Y is not substituted or is substituted by at least one halogen group (preferably fluoro or chloro).

For example, preferred compounds of the invention include those wherein B is phenyl, pyridyl, pyrimidinyl, or thienyl.

Other preferred compounds include those wherein  $L_1$  is a direct bond,  $L_2$  is a direct bond,  $M_1$  is N,  $M_2$  is C or N,  $M_3$  is N,  $M_4$  is C, and  $M_5$  is C.

Other preferred compounds include those wherein R1 is 3- or 4- chlorophenylethyl, 3- or 4- chlorophenylpropyl, 2- or 3-pyridylethyl, 2- or 3-pyridylpropyl, 2- or 4-pyrimidinylethyl (optionally monosubstituted by fluoro or chloro), 2- or 4-pyrimidinylpropyl (optionally monosubstituted by fluoro or chloro), 2-(2-pyrimidinyl)ethyl (optionally monosubstituted by fluoro or chloro), 2-(2-pyrimidinyl)propyl (optionally monosubstituted by fluoro or chloro). Particularly preferred compounds include those wherein R1 is 2-pyrimidinylpropyl, 2-pyrimidinylethyl or 5-fluoro-2-pyrimidinylethyl.

It will be appreciated that the particular substituents and number of substituents on B and/or R1 are selected so as to avoid sterically undesirable combinations.

Each exemplified compound represents a particular and independent aspect of the invention.

Where optically active centres exist in the compounds of formulae I or II, we disclose all individual optically active forms and combinations of these as individual specific embodiments of the invention, as well as their corresponding racemates.

It will be appreciated that the compounds according to the invention can contain one or more asymmetrically substituted carbon atoms. The presence of one or more of these asymmetric centres (chiral centres) in a compound of formulae I or II can give rise to stereoisomers, and in each case the invention is to be understood to extend to all such stereoisomers, including enantiomers and diastereomers, and mixtures including racemic mixtures thereof.

Where tautomers exist in the compounds of formulae I or II, we disclose all individual tautomeric forms and combinations of these as individual specific embodiments of the invention.

As previously outlined the compounds of the invention are metalloproteinase inhibitors, in particular they are inhibitors of MMP13. Each of the above indications for the compounds of the formulae I or II represents an independent and particular embodiment of the invention. Whilst we do not wish to be bound by theoretical considerations, the compounds of the invention are believed to show selective inhibition for any one of the above indications relative to any MMP1 inhibitory activity, by way of non-limiting example they may show 100-1000 fold selectivity over any MMP1 inhibitory activity.

The compounds of the invention may be provided as pharmaceutically acceptable salts. These include acid addition salts such as hydrochloride, hydrobromide, citrate and maleate salts and salts formed with phosphoric and sulphuric acid. In another aspect suitable salts are base salts such as an alkali metal salt for example sodium or potassium, an alkaline earth metal salt for example calcium or magnesium, or organic amine salt for example triethylamine.

They may also be provided as *in vivo* hydrolysable esters. These are pharmaceutically acceptable esters that hydrolyse in the human body to produce the parent compound. Such esters can be identified by administering, for example intravenously to a test animal, the compound under test and subsequently examining the test animal's body fluids. Suitable *in vivo* hydrolysable esters for carboxy include methoxymethyl and for hydroxy include formyl and acetyl, especially acetyl.

In order to use a compound of the formulae I or II or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof for the therapeutic treatment (including



prophylactic treatment) of mammals including humans, it is normally formulated in accordance with standard pharmaceutical practice as a pharmaceutical composition.

Therefore in another aspect the present invention provides a pharmaceutical composition which comprises a compound of the formulae I or II or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester and pharmaceutically acceptable carrier.

The pharmaceutical compositions of this invention may be administered in standard manner for the disease or condition that it is desired to treat, for example by oral, topical, parenteral, buccal, nasal, vaginal or rectal administration or by inhalation. For these purposes the compounds of this invention may be formulated by means known in the art into the form of, for example, tablets, capsules, aqueous or oily solutions, suspensions, emulsions, creams, ointments, gels, nasal sprays, suppositories, finely divided powders or aerosols for inhalation, and for parenteral use (including intravenous, intramuscular or infusion) sterile aqueous or oily solutions or suspensions or sterile emulsions.

In addition to the compounds of the present invention the pharmaceutical composition of this invention may also contain, or be co-administered (simultaneously or sequentially) with, one or more pharmacological agents of value in treating one or more diseases or conditions referred to hereinabove.

The pharmaceutical compositions of this invention will normally be administered to humans so that, for example, a daily dose of 0.5 to 75 mg/kg body weight (and preferably of 0.5 to 30 mg/kg body weight) is received. This daily dose may be given in divided doses as necessary, the precise amount of the compound received and the route of administration depending on the weight, age and sex of the patient being treated and on the particular disease or condition being treated according to principles known in the art.

Typically unit dosage forms will contain about 1 mg to 500 mg of a compound of this invention.

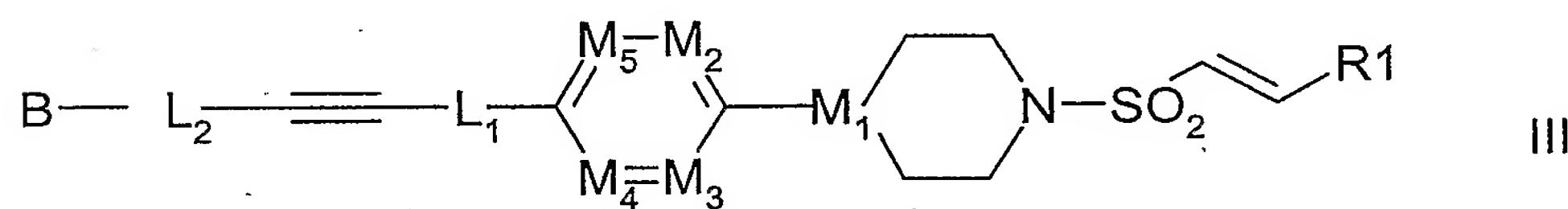
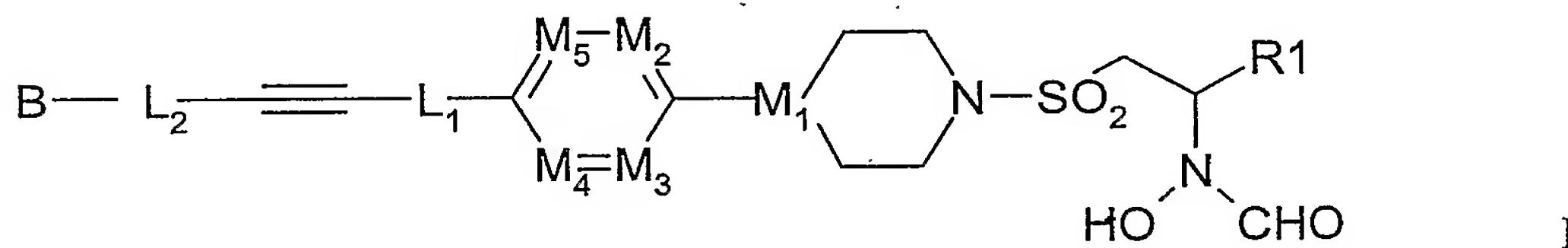
Therefore in a further aspect, the present invention provides a compound of the formulae I or II or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof for use in a method of therapeutic treatment of the human or animal body. In particular we disclose use in the treatment of a disease or condition mediated by MMP13.

In yet a further aspect the present invention provides a method of treating a metalloproteinase mediated disease or condition which comprises administering to a warm-

blooded animal a therapeutically effective amount of a compound of the formulae I or II or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof. Metalloproteinase mediated diseases or conditions include arthritis (such as osteoarthritis), atherosclerosis, chronic obstructive pulmonary diseases (COPD).

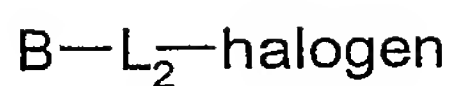
In another aspect the present invention provides a process for preparing a compound of the formulae I or II or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof which process is outlined below.

A compound of the formula I can be prepared from a compound of the formula III by reaction with hydroxylamine followed by formylation.

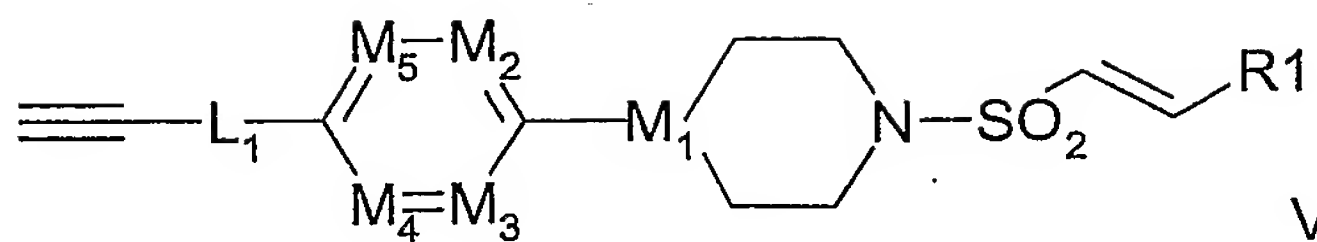


A compound of the formula III can be prepared from a compound of the formula IV and a compound of the formula V or from a compound of the formula VI and a compound of the formula VII. A compound of the formula V may be prepared from a compound of the formula VII.

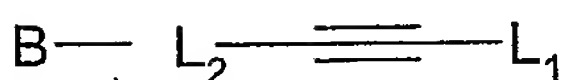
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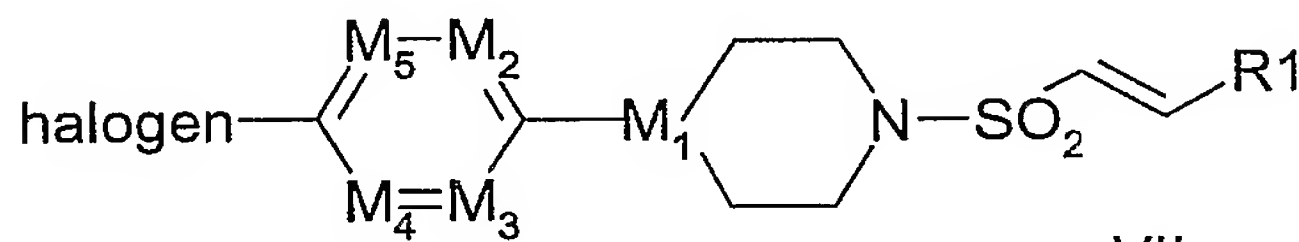
IV



V

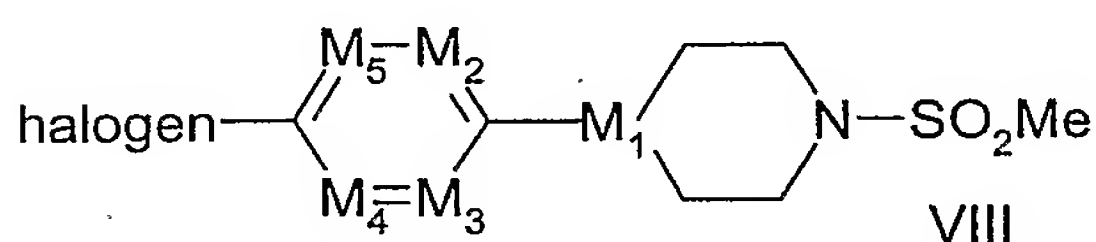


VI

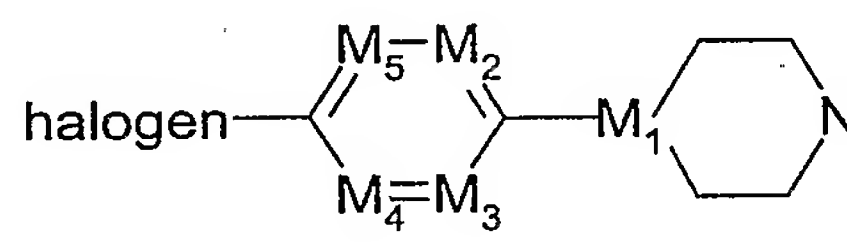


VII

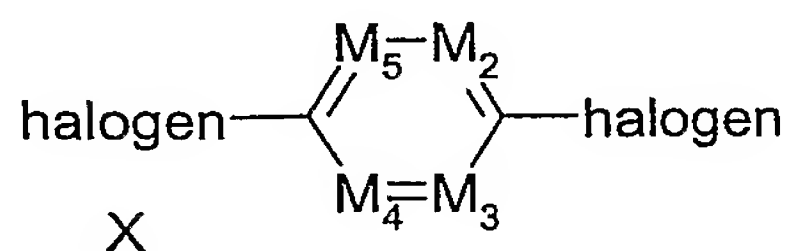
A compound of the formula VII may be prepared from a compound of the formula VIII by reaction with a suitable aldehyde (R1CHO) or a suitable ester (R1COOR). A compound of formula VIII may be conveniently prepared from a compound of formula IX. A compound of the formula IX can be prepared conveniently from a compound of the formula X and a compound of the formula XI (where P is hydrogen or a suitable protecting group and M<sub>1</sub>' is hydrogen or a suitably reactive group).



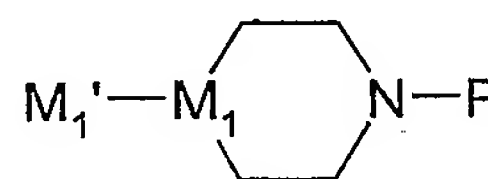
VIII



IX



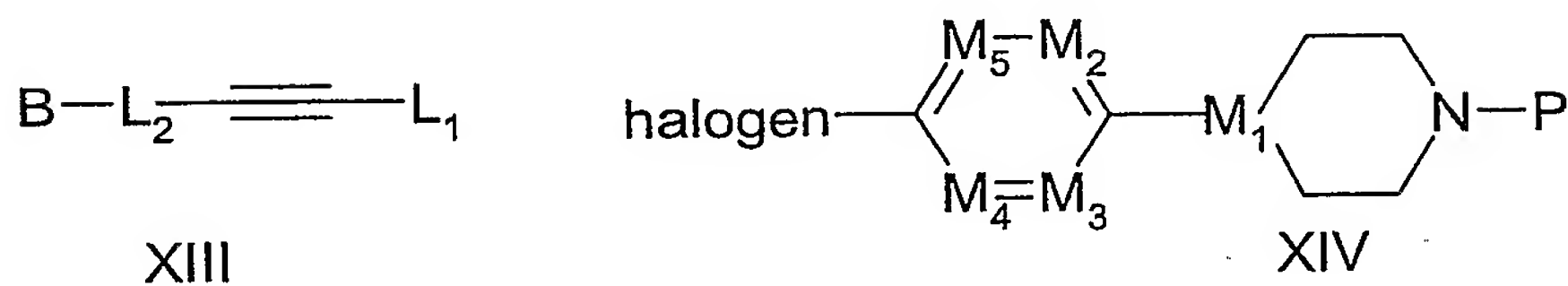
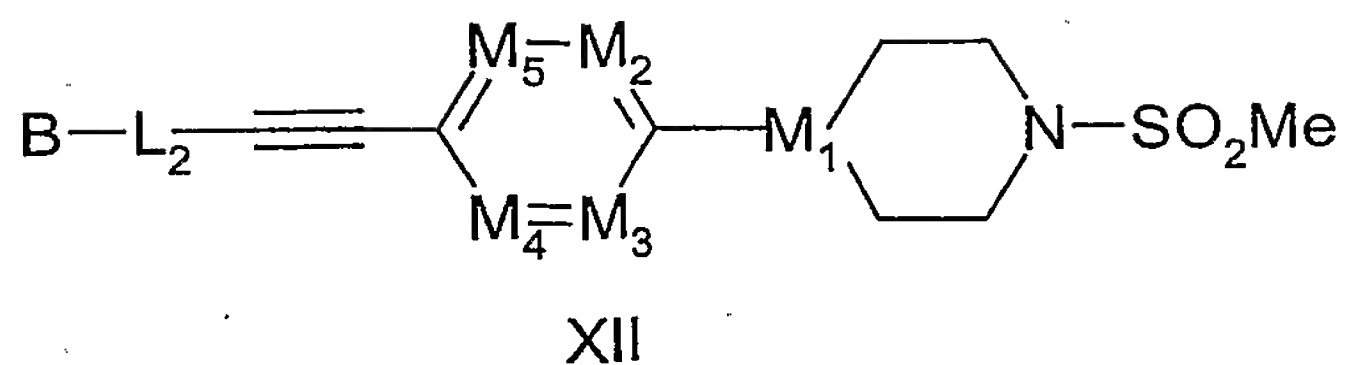
X



XI

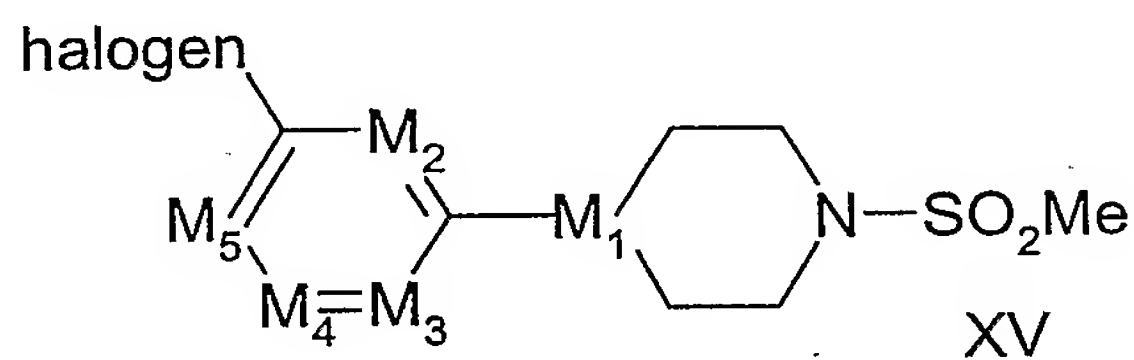
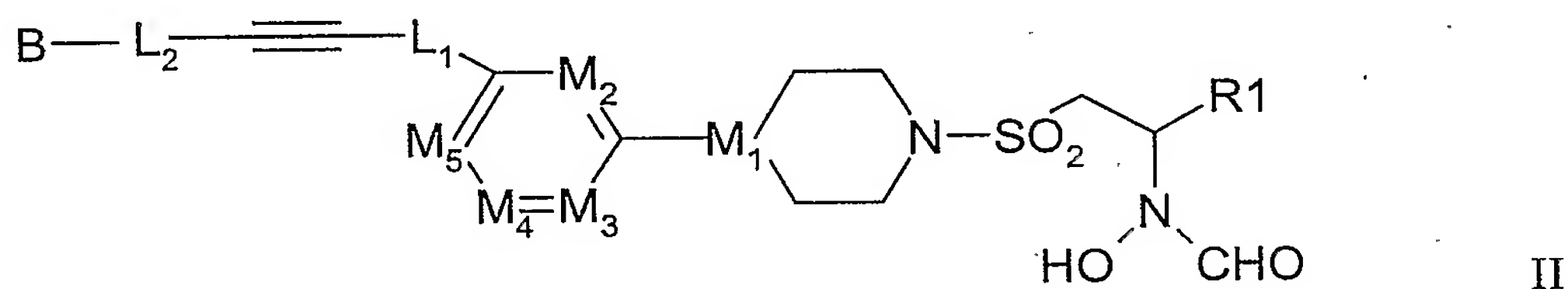
A compound of the formula III can also be prepared from a compound of the formula XII by reaction with a suitable aldehyde (R1CHO) or a suitable ester (R1COOR). A compound of the formula XII can be prepared conveniently from a compound of the formula XIII and a compound of the formula XIV

11



A compound of the formula II may be prepared from a compound of the formula XV by similar methodology to that described for compound I above.

5



10 It will be appreciated that many of the relevant starting materials are commercially available or may be made by any convenient method as described in the literature or known to the skilled chemist or described in the Examples herein.

15

The compounds of the invention may be evaluated for example in the following assays:

## Isolated Enzyme Assays

### Matrix Metalloproteinase family including for example MMP13.

Recombinant human proMMP13 may be expressed and purified as described by Knauper *et al.* [V. Knauper *et al.*, (1996) The Biochemical Journal 271:1544-1550 (1996)]. The purified enzyme can be used to monitor inhibitors of activity as follows: purified proMMP13 is activated using 1mM amino phenyl mercuric acid (APMA), 20 hours at 21°C; the activated MMP13 (11.25ng per assay) is incubated for 4-5 hours at 35°C in assay buffer (0.1M Tris-HCl, pH 7.5 containing 0.1M NaCl, 20mM CaCl<sub>2</sub>, 0.02 mM ZnCl<sub>2</sub> and 0.05% (w/v) Brij 35 using the synthetic substrate 7-methoxycoumarin-4-yl)acetyl.Pro.Leu.Gly.Leu.N-3-(2,4-dinitrophenyl)-L-2,3-diaminopropionyl.Ala.Arg.NH<sub>2</sub> in the presence or absence of inhibitors. Activity is determined by measuring the fluorescence at  $\lambda_{ex}$  328nm and  $\lambda_{em}$  393nm. Percent inhibition is calculated as follows: % Inhibition is equal to the [Fluorescence<sub>plus inhibitor</sub> - Fluorescence<sub>background</sub>] divided by the [Fluorescence<sub>minus inhibitor</sub> - Fluorescence<sub>background</sub>].

A similar protocol can be used for other expressed and purified pro MMPs using substrates and buffers conditions optimal for the particular MMP, for instance as described in C. Graham Knight *et al.*, (1992) FEBS Lett. 296(3):263-266.

### Adamalysin family including for example TNF convertase

The ability of the compounds to inhibit proTNF $\alpha$  convertase enzyme may be assessed using a partially purified, isolated enzyme assay, the enzyme being obtained from the membranes of THP-1 as described by K. M. Mohler *et al.*, (1994) Nature 370:218-220. The purified enzyme activity and inhibition thereof is determined by incubating the partially purified enzyme in the presence or absence of test compounds using the substrate 4',5'-Dimethoxy-fluoresceinyl Ser.Pro.Leu.Alu.Gln.Alu.Val.Arg.Ser.Ser.Ser.Arg.Cys(4-(3-succinimid-1-yl)-fluorescein)-NH<sub>2</sub> in assay buffer (50mM Tris HCl, pH 7.4 containing 0.1% (w/v) Triton X-100 and 2mM CaCl<sub>2</sub>), at 26°C for 18 hours. The amount of inhibition is determined as for MMP13 except  $\lambda_{ex}$  490nm and  $\lambda_{em}$  530nm were used. The substrate was synthesised as follows. The peptidic part of the substrate was assembled on Fmoc-NH-Rink-MBHA-polystyrene resin either manually or on an automated peptide synthesiser



by standard methods involving the use of Fmoc-amino acids and O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate (HBTU) as coupling agent with at least a 4- or 5-fold excess of Fmoc-amino acid and HBTU. Ser<sup>1</sup> and Pro<sup>2</sup> were double-coupled. The following side chain protection strategy was employed; Ser<sup>1</sup>(But), Gln<sup>5</sup>(Trityl), Arg<sup>8,12</sup>(Pmc or Pbf), Ser<sup>9,10,11</sup>(Trityl), Cys<sup>13</sup>(Trityl). Following assembly, the N-terminal Fmoc-protecting group was removed by treating the Fmoc-peptidyl-resin with 20% piperidine in DMF. The amino-peptidyl-resin so obtained was acylated by treatment for 1.5-2hr at 70°C with 1.5-2 equivalents of 4',5'-dimethoxy-fluorescein-4(5)-carboxylic acid [Khanna & Ullman, (1980) Anal Biochem. 108:156-161] which had been preactivated with diisopropylcarbodiimide and 1-hydroxybenzotriazole in DMF]. The dimethoxyfluoresceinyl-peptide was then simultaneously deprotected and cleaved from the resin by treatment with trifluoroacetic acid containing 5% each of water and triethylsilane. The dimethoxyfluoresceinyl-peptide was isolated by evaporation, trituration with diethyl ether and filtration. The isolated peptide was reacted with 4-(N-maleimido)-fluorescein in DMF containing diisopropylethylamine, the product purified by RP-HPLC and finally isolated by freeze-drying from aqueous acetic acid. The product was characterised by MALDI-TOF MS and amino acid analysis.

### Natural Substrates

The activity of the compounds of the invention as inhibitors of aggrecan degradation may be assayed using methods for example based on the disclosures of E. C. Arner *et al.*, (1998) Osteoarthritis and Cartilage 6:214-228; (1999) Journal of Biological Chemistry, 274 (10), 6594-6601 and the antibodies described therein. The potency of compounds to act as inhibitors against collagenases can be determined as described by T. Cawston and A. Barrett (1979) Anal. Biochem. 99:340-345.

### Inhibition of metalloproteinase activity in cell/tissue based activity

#### Test as an agent to inhibit membrane sheddases such as TNF convertase

The ability of the compounds of this invention to inhibit the cellular processing of TNF $\alpha$  production may be assessed in THP-1 cells using an ELISA to detect released TNF essentially as described K. M. Mohler *et al.*, (1994) Nature 370:218-220. In a similar

fashion the processing or shedding of other membrane molecules such as those described in N. M. Hooper *et al.*, (1997) *Biochem. J.* 321:265-279 may be tested using appropriate cell lines and with suitable antibodies to detect the shed protein.

5    **Test as an agent to inhibit cell based invasion**

The ability of the compound of this invention to inhibit the migration of cells in an invasion assay may be determined as described in A. Albini *et al.*, (1987) *Cancer Research* 47:3239-3245.

10   **Test as an agent to inhibit whole blood TNF sheddase activity**

The ability of the compounds of this invention to inhibit TNF $\alpha$  production is assessed in a human whole blood assay where LPS is used to stimulate the release of TNF $\alpha$ . Heparinized (10Units/mL) human blood obtained from volunteers is diluted 1:5 with medium (RPMI1640 + bicarbonate, penicillin, streptomycin and glutamine) and incubated  
15 (160 $\mu$ l) with 20 $\mu$ l of test compound (triplicates), in DMSO or appropriate vehicle, for 30 min at 37°C in a humidified (5%CO<sub>2</sub>/95%air) incubator, prior to addition of 20 $\mu$ l LPS (*E. coli*. 0111:B4; final concentration 10 $\mu$ g/mL). Each assay includes controls of diluted blood incubated with medium alone (6 wells/plate) or a known TNF $\alpha$  inhibitor as standard. The plates are then incubated for 6 hours at 37°C (humidified incubator), centrifuged  
20 (2000rpm for 10 min; 4°C ), plasma harvested (50-100 $\mu$ l) and stored in 96 well plates at -70°C before subsequent analysis for TNF $\alpha$  concentration by ELISA.

**Test as an agent to inhibit in vitro cartilage degradation**

The ability of the compounds of this invention to inhibit the degradation of the  
25 aggrecan or collagen components of cartilage can be assessed essentially as described by K. M. Bottomley *et al.*, (1997) *Biochem J.* 323:483-488.

**Pharmacodynamic test**

To evaluate the clearance properties and bioavailability of the compounds of this  
30 invention an ex vivo pharmacodynamic test is employed which utilises the synthetic substrate assays above or alternatively HPLC or Mass spectrometric analysis. This is a

generic test which can be used to estimate the clearance rate of compounds across a range of species. Animals (e.g. rats, marmosets) are dosed iv or po with a soluble formulation of compound (such as 20% w/v DMSO, 60% w/v PEG400) and at subsequent time points (e.g. 5, 15, 30, 60, 120, 240, 480, 720, 1220 mins) the blood samples are taken from an appropriate vessel into 10U heparin. Plasma fractions are obtained following centrifugation and the plasma proteins precipitated with acetonitrile (80% w/v final concentration). After 30 mins at -20°C the plasma proteins are sedimented by centrifugation and the supernatant fraction is evaporated to dryness using a Savant speed vac. The sediment is reconstituted in assay buffer and subsequently analysed for compound content using the synthetic substrate assay. Briefly, a compound concentration-response curve is constructed for the compound undergoing evaluation. Serial dilutions of the reconstituted plasma extracts are assessed for activity and the amount of compound present in the original plasma sample is calculated using the concentration-response curve taking into account the total plasma dilution factor.

### **In vivo assessment**

#### **Test as an anti-TNF agent**

The ability of the compounds of this invention as *ex vivo* TNF $\alpha$  inhibitors is assessed in the rat. Briefly, groups of male Wistar Alderley Park (AP) rats (180-210g) are dosed with compound (6 rats) or drug vehicle (10 rats) by the appropriate route e.g. peroral (p.o.), intraperitoneal (i.p.), subcutaneous (s.c.). Ninety minutes later rats are sacrificed using a rising concentration of CO<sub>2</sub> and bled out via the posterior vena cavae into 5 Units of sodium heparin/mL blood. Blood samples are immediately placed on ice and centrifuged at 2000 rpm for 10 min at 4°C and the harvested plasmas frozen at -20°C for subsequent assay of their effect on TNF $\alpha$  production by LPS-stimulated human blood. The rat plasma samples are thawed and 175 $\mu$ l of each sample are added to a set format pattern in a 96 well plate. Fifty  $\mu$ l of heparinized human blood is then added to each well, mixed and the plate is incubated for 30 min at 37°C (humidified incubator). LPS (25 $\mu$ l; final concentration 10 $\mu$ g/mL) is added to the wells and incubation continued for a further 5.5 hours. Control wells are incubated with 25 $\mu$ l of medium alone. Plates are then centrifuged

for 10 min at 2000 rpm and 200µl of the supernatants are transferred to a 96 well plate and frozen at -20°C for subsequent analysis of TNF concentration by ELISA.

Data analysis by dedicated software calculates for each compound/dose:

5    Percent inhibition of TNFα = 
$$\frac{\text{Mean TNF}\alpha \text{ (Controls)} - \text{Mean TNF}\alpha \text{ (Treated)}}{\text{Mean TNF}\alpha \text{ (Controls)}} \times 100$$

**Test as an anti-arthritic agent**

10    Activity of a compound as an anti-arthritic is tested in the collagen-induced arthritis (CIA) as defined by D. E. Trentham *et al.*, (1977) J. Exp. Med. 146:857. In this model acid soluble native type II collagen causes polyarthritis in rats when administered in Freund's incomplete adjuvant. Similar conditions can be used to induce arthritis in mice and primates.

15    **Test as an anti-cancer agent**

Activity of a compound as an anti-cancer agent may be assessed essentially as described in I. J. Fidler (1978) Methods in Cancer Research 15:399-439, using for example the B16 cell line (described in B. Hibner *et al.*, Abstract 283 p75 10th NCI-EORTC Symposium, Amsterdam June 16 – 19 1998).

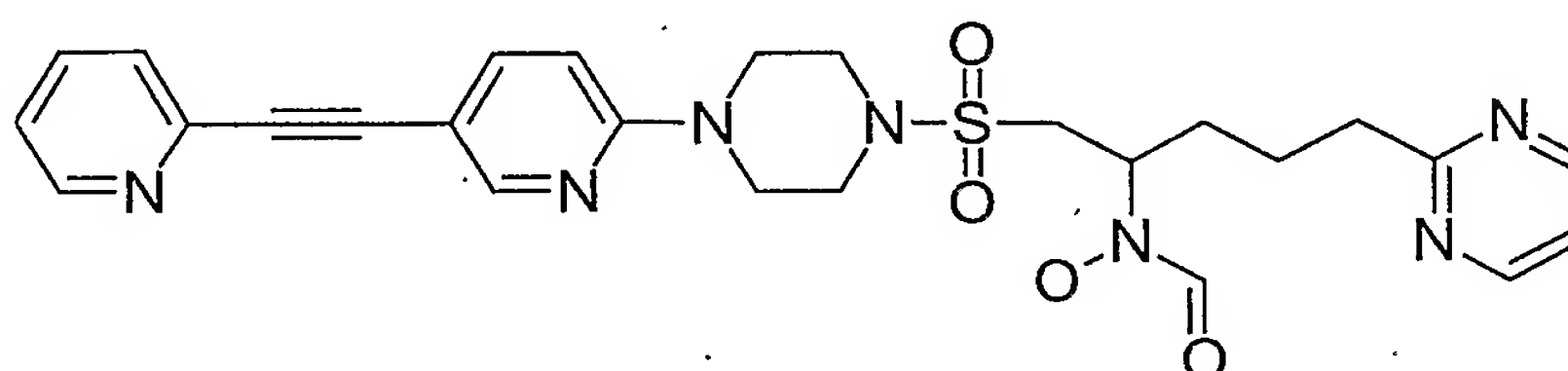
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The invention will now be illustrated but not limited by the following Examples:

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**EXAMPLE 1**

**hydroxy{-1-[(4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl}formamide**



To formic acid (1.5 mL, 39 mmol) at 0 °C was added acetic anhydride (375 µL, 3.9 mmol) and the mixture was stirred at RT for 10 minutes. The reaction was then recooled to 0 °C, and a solution of 2-[4-(hydroxyamino)-5-({4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pentyl]pyrimidine (403 mg, 0.79 mmol) in THF (8 mL) was then added. The reaction was brought to RT and stirred for one hour. Volatiles were then removed *in vacuo*, and the residue azeotroped with toluene (2 x 5 mL). The residue was then dissolved in MeOH (10 mL) and heated to 40 °C for one hour. The solution was then cooled to RT and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 10% MeOH in EtOAc) to give the title compound as a pale yellow foam (231 mg, 0.43 mmol, 54%).

<sup>1</sup>H NMR (DMSO-D<sub>6</sub>, 373K) : 9.41 (br s, 1 H), 8.65 (d, 2H), 8.56 (d, 1H), 8.36 (d, 1 H), 8.11 (br s, 1 H), 7.79 (m, 1 H), 7.71 (dd, 1 H), 7.55 (d, 1 H), 7.32 (m, 1 H), 7.28 (dd, 1 H), 6.88 (d, 1 H), 3.72 (m, 4 H), 3.45 (dd, 1 H), 3.32 (m, 4 H), 3.17 (dd, 1 H), 2.92 (m, 3 H), 1.77 (m, 4 H)

MS (ESI): 536.45 (MH<sup>+</sup>)

The starting material was prepared as follows:

To a stirred solution of 2-chloro-5-iodo-pyridine (CAS number 69045-79-0, 10.52 g, 43.9 mmol) and diisopropylethylamine (11.5 mL, 65.9 mmol) in DMA (200 mL) was added piperazine (15.14 g, 0.176 mol). The mixture was then heated to 120°C for 20 hours. The solid precipitate was then filtered off, and the filtrate evaporated *in vacuo*. The residue



was partitioned between EtOAc (100 mL) and water (100 mL) and the layers were separated. The aqueous layer was then extracted with EtOAc (2 x 100 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*, to give 1-(5-iodopyridin-2-yl)piperazine as a yellow solid (10.42 g, 36 mmol, 82%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.30 (d, 1 H), 7.65 (dd, 1 H), 6.45 (d, 1 H), 3.46, (t, 4 H), 2.97 (t, 4 H).  
MS (ESI): 290.29 (MH<sup>+</sup>)

To a stirred solution of 1-(5-iodopyridin-2-yl)piperazine (CAS number 219635-89-9, 10.42 g, 36 mmol), in CH<sub>2</sub>Cl<sub>2</sub> (150 mL) at 0°C was added a solution of BOC-O-BOC (7.87 g, 26 mmol). The reaction was allowed to warm to RT and was stirred for 18 hours. Volatiles were removed *in vacuo*, and the residue was dissolved in EtOAc (200 mL). The solution was washed with water (100 mL), and the aqueous layer was extracted with EtOAc (2 x 100 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*, to give *tert*-butyl 4-(5-iodopyridin-2-yl)piperazine-1-carboxylate as a yellow solid (14 g, 36 mmol, 99%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.31 (d, 1 H), 7.08 (dd, 1 H), 6.46 (d, 1 H), 3.50 (s, 8H), 1.48 (s, 9 H).  
MS (ESI): 390.37 (MH<sup>+</sup>)

To a stirred solution of *tert*-butyl 4-(5-iodopyridin-2-yl)piperazine-1-carboxylate (4.087 g, 10.5 mmol) in DMA (75 mL) at RT was added 2-ethynylpyridine, (CAS number 1945-84-2, 1.17 mL, 11.5 mmol), triethylamine (4.4 mL, 31.5 mmol), CuI (800 mg, 4.2 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (1.21 g, 1.05 mmol). After stirring for 30 minutes, the DMA was removed *in vacuo*, and the residue diluted with EtOAc (100 mL). The dark solution was then washed with water (100 mL). The layers were separated and the aqueous phase extracted with EtOAc (3 x 50 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 50% EtOAc in hexanes) to give *tert*-butyl 4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazine-1-carboxylate as a brown oil (3.8 g, 10.5 mmol, 99%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.62 (m, 1 H), 8.43 (d, 1 H), 7.68 (m, 2 H), 7.49 (d, 1 H), 7.21 (dd, 1 H), 6.58 (d, 1 H), 3.61 (m, 8 H), 1.51 (s, 9 H).

MS (ESI): 364.94 (MH<sup>+</sup>)

5 To a stirred solution of *tert*-butyl 4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazine-1-carboxylate (3.8 g, 10.4 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (35 mL) at 0°C was added trifluoroacetic acid (17.2 mL). The reaction was brought to room temperature and stirred for 30 minutes. Volatiles were removed *in vacuo*, and the residue was azeotroped with toluene (2 x 10 mL). The residue was then dissolved in CH<sub>2</sub>Cl<sub>2</sub> and washed with aqueous sodium  
10 hydroxide solution (2M, 2 x 20 mL). The layers were then separated and the aqueous phase extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 x 10 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (50 mL), and cooled to 0°C and treated sequentially with triethylamine (1.5 mL, 16.7 mmol) and methanesulfonyl chloride (1.1 mL, 13.5 mmol). The reaction was brought to RT and  
15 stirred for 1 hour. The reaction was then quenched with water (50 mL). The layers were separated and the aqueous phase extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 x 20 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 10% MeOH in EtoAc) to give 1-(methanesulfonyl)-4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazine as a brown solid (3.5 g,  
20 10.2 mmol, 98%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.60 (d, 1 H), 8.46 (d, 1 H), 7.69, (m, 2 H), 7.52 (d, 1 H), 7.23 (m, 1 H), 6.65 (d, 1 H), 3.79 (m, 4 H), 3.35 (m, 4 H), 2.83 (s, 3 H).

MS (ESI): 342.83 (MH<sup>+</sup>)

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To a stirred suspension of 1-(methanesulfonyl)-4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazine (456 mg, 1.33 mmol) in THF (10 mL) at -20°C, was added dropwise a solution of LiHMDS in THF (3.0 mL, 1.0M solution, 3.0 mmol). The resulting suspension was stirred at -20°C for 30 minutes before being treated with diethyl chlorophosphate  
30 (212 µL, 1.46 mmol). The solution was then maintained at -20°C for 15 minutes before being treated with a solution of 4-pyrimidin-2-ylbutanal (220 mg, 1.46 mmol, CAS number

260441-10-9) in THF (3 mL). The solution was stirred at  $-20^{\circ}\text{C}$  for a further 20 minutes before being quenched with saturated aqueous ammonium chloride solution (10 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 10 mL). The combined organic extracts were then dried, ( $\text{MgSO}_4$ ), filtered and concentrated *in vacuo* to give -[(4*E/Z*)-5-({4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pent-4-enyl]pyrimidine. This material was used crude in the next step.

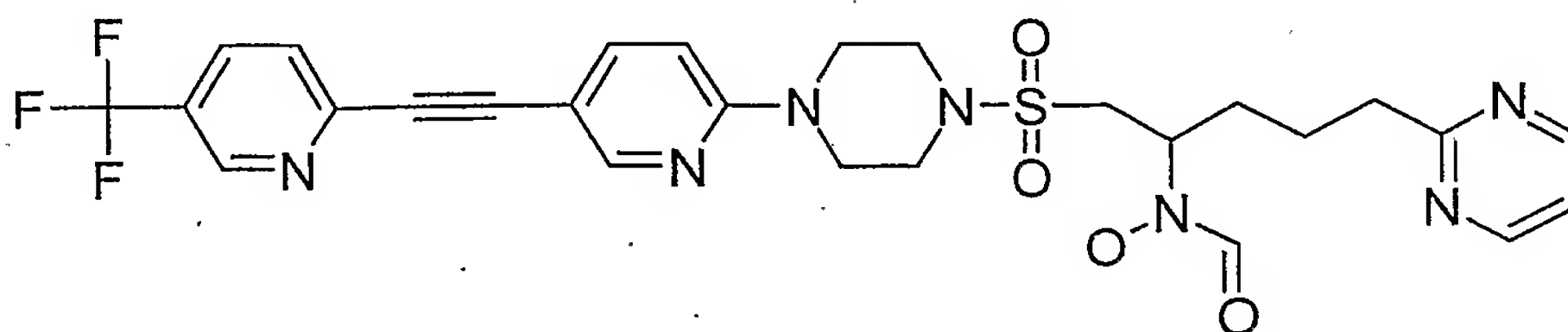
MS (ESI): 474.97 ( $\text{MH}^+$ )

To a stirred solution of 2-[(4*E/Z*)-5-({4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pent-4-enyl]pyrimidine (crude from previous step), in THF (15 mL) at RT was added a solution of hydroxylamine (3 mL, 50% aqueous solution in water). The reaction was stirred for 3 hours at RT before being quenched with saturated aqueous ammonium chloride solution (5 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 10 mL). The combined organic extracts were then dried, ( $\text{MgSO}_4$ ), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica, 10% MeOH in ethyl acetate) to give 2-[4-(hydroxyamino)-5-({4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pentyl]pyrimidine (403 mg, 0.79 mmol).

MS (ESI): 508.22 ( $\text{MH}^+$ )

## EXAMPLE 2

hydroxy[4-pyrimidin-2-yl-1-({[4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazin-1-yl]sulfonyl}methyl)butyl]formamide



To formic acid (0.36 mL, 9.5 mmol) at 0 °C was added acetic anhydride (90 µL, 0.95 mmol) and the mixture was stirred at RT for 10 minutes. The reaction was then recooled to 0 °C, and added to a solution of 2-(4-(hydroxyamino)-5-{{4-(5-{{5-(trifluoromethyl)pyridin-2-yl}ethynyl}pyridin-2-yl)piperazin-1-yl}sulfonyl}pentyl)pyrimidine (403 mg, 0.79 mmol) and formic acid (0.36 mL, 9.5 mmol) in THF (3 mL) at 0 °C. The reaction was brought to RT and stirred for one hour. Volatiles were then removed *in vacuo*, and the residue azeotroped with toluene (2 x 5 mL). The residue was then dissolved in MeOH (10 mL) and heated to 40 °C for one hour. The solution was then cooled to RT and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 10% MeOH in EtOAc) to give the title compound as a pale yellow foam (66 mg, 0.11 mmol, 58%).

<sup>1</sup>H NMR (DMSO-D<sub>6</sub>, 373K) : 9.38 (br s, 1 H), 8.90 (d, 1 H), 8.67 (d, 2 H), 8.42 (d, 1 H), 8.15 (dd, 1 H), 8.14 (br s, 1 H), 7.73 (m, 2 H), 7.27 (t, 1 H), 6.90 (d, 1 H), 3.71 (m, 4 H), 3.42 (dd, 1 H), 3.30 (m, 4 H), 3.16 (dd, 1 H), 2.88 (m, 3 H), 1.76 (m, 3 H), 1.66 (m, 1H)  
MS (ESI): 604.39 (MH<sup>+</sup>)

The starting material was prepared as follows:

To a stirred solution of *tert*-butyl 4-(5-iodopyridin-2-yl)piperazine-1-carboxylate (1.0 g, 2.57 mmol, prepared as in Example 1), trimethylsilyl acetylene (0.73 mL, 5.14 mmol), triethylamine (1.11 mL, 7.71 mmol), and cuprous iodide (196 mg, 1.03 mmol) in DMA (20 mL) at 25 °C was added tetrakis(triphenylphosphine) palladium (0) (149 mg, 5mol%). The reaction was stirred for 10 minutes. Volatiles were removed *in vacuo*. The residue was then dissolved in ethyl acetate (250 mL) and washed with water (100 mL). The layers were then separated and the aqueous phase extracted with ethyl acetate (250 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 10% EtOAc in hexanes) to give *tert*-butyl 4-{5-[(trimethylsilyl)ethynyl]pyridin-2-yl}piperazine-1-carboxylate as a pale brown solid (652 mg, 1.81 mmol, 70%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.28 (d, 1 H), 7.50 (dd, 1 H), 6.50 (d, 1 H), 3.57 (m, 8 H), 1.49 (s, 9 H), 0.25 (s, 9H).

MS (ESI): 360.51 (MH<sup>+</sup>)

5 To a stirred solution of *tert*-butyl 4-{5-[(trimethylsilyl)ethynyl]pyridin-2-yl}piperazine-1-carboxylate (637 mg, 1.77 mmol), in THF (8 mL) at 25<sup>0</sup>C was added tetrabutylammonium fluoride (1.77 mL, 1.0M in THF). The reaction was stirred for 1 hour. Water (5 mL) was then added and product extracted with ethyl acetate (2 x 5 mL). The combined organic  
10 extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo* to give *tert*-butyl 4-(5-ethynylpyridin-2-yl)piperazine-1-carboxylate as an orange solid (509 mg, 1.77 mmol, 100%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.30 (d, 1 H), 7.54 (dd, 1 H), 6.52 (d, 1 H), 3.56 (m, 8H), 3.07 (s, 1 H), 1.49 (s, 9H).

15 To a stirred solution of *tert*-butyl 4-(5-ethynylpyridin-2-yl)piperazine-1-carboxylate (509 mg, 1.77 mmol), 2-Bromo-5-trifluoromethyl pyridine (400mg, 1.77 mmol), triethylamine (0.74 mL, 5.31 mmol), and cuprous iodide (135mg, 0.71 mmol) in DMA (20 mL) at 25<sup>0</sup>C was added tetrakis(triphenylphosphine) palladium (0) ( 102 mg, 5 mol%). The reaction was then stirred for 1 hour. Volatiles were removed *in vacuo*, and the residue was azeotroped  
20 with toluene (2 x 10 mL). The residue was then dissolved in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and washed with water (10 mL). The layers were then separated and the aqueous phase extracted with CH<sub>2</sub>Cl<sub>2</sub> (2x 100 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 25% EtOAc in hexanes) to give *tert*-butyl 4-(5-{[5-(trifluoromethyl)pyridin-2-  
25 yl]ethynyl}pyridin-2-yl)piperazine-1-carboxylate as a yellow solid (729 mg, 1.68 mmol, 95%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.83 (d, 1 H), 8.42 (d, 1 H), 7.88, (dd, 1 H), 7.67 (dd, 1 H), 7.56 (d, 1 H), 6.59 (d, 1 H), 3.60 (m, 4 H), 3.53 (m, 4 H), 1.50 (s, 9H).

30 MS (ESI): 433.49 (MH<sup>+</sup>)



To a stirred solution of *tert*-butyl 4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazine-1-carboxylate (724 mg, 1.67 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (8 mL) at 0°C was added trifluoroacetic acid (4 mL). The reaction was brought to room temperature and stirred for 30 minutes. Volatiles were removed *in vacuo*. The residue was then dissolved in CH<sub>2</sub>Cl<sub>2</sub> (20 mL) and washed with aqueous sodium hydroxide solution (2M, 2 x 20 mL). The layers were then separated and the aqueous phase extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 x 10 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (50 mL), and cooled to 0°C and treated sequentially with triethylamine (0.25 mL, 1.75 mmol) and methanesulfonyl chloride (0.14 mL, 13.5 mmol). The reaction was brought to RT and stirred for 1 hour. The reaction was then quenched with water (50 mL). The layers were separated and the aqueous phase extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 x 20 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo* to give 1-(methylsulfonyl)-4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazine as an orange solid (445 mg, 1.08 mmol, 65%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>): 8.85 (d, 1 H), 8.45 (d, 1 H), 7.90, (dd, 1 H), 7.70 (dd, 1 H), 7.60 (d, 1 H), 6.65 (d, 1 H), 3.79 (m, 4 H), 3.35 (t, 4 H), 2.80 (s, 3 H).

MS (ESI): 411.12 (MH<sup>+</sup>)

To a stirred suspension of 1-(methylsulfonyl)-4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazine (437 mg, 1.06 mmol) in THF (10 mL) at -20°C, was added dropwise a solution of LiHMDS in THF (2.34 mL, 1.0M solution, 2.34 mmol). The resulting suspension was stirred at -20°C for 30 minutes before being treated with diethyl chlorophosphate (0.17 mL, 1.17 mmol). The solution was then maintained at -20°C for 15 minutes before being treated with a solution of 4-pyrimidin-2-ylbutanal (159 mg, 1.06 mmol, CAS number 260441-10-9) in THF (3 mL). The solution was stirred at -20°C for a further 20 minutes before being quenched with saturated aqueous ammonium chloride solution (10 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 10 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo* to give 2-((4*E/Z*)-5-{[4-(5-{[5-(trifluoromethyl)pyridin-2-

yl]ethynyl}pyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine. This was used crude in the next step.

MS (ESI): 543.19 (MH<sup>+</sup>)

5

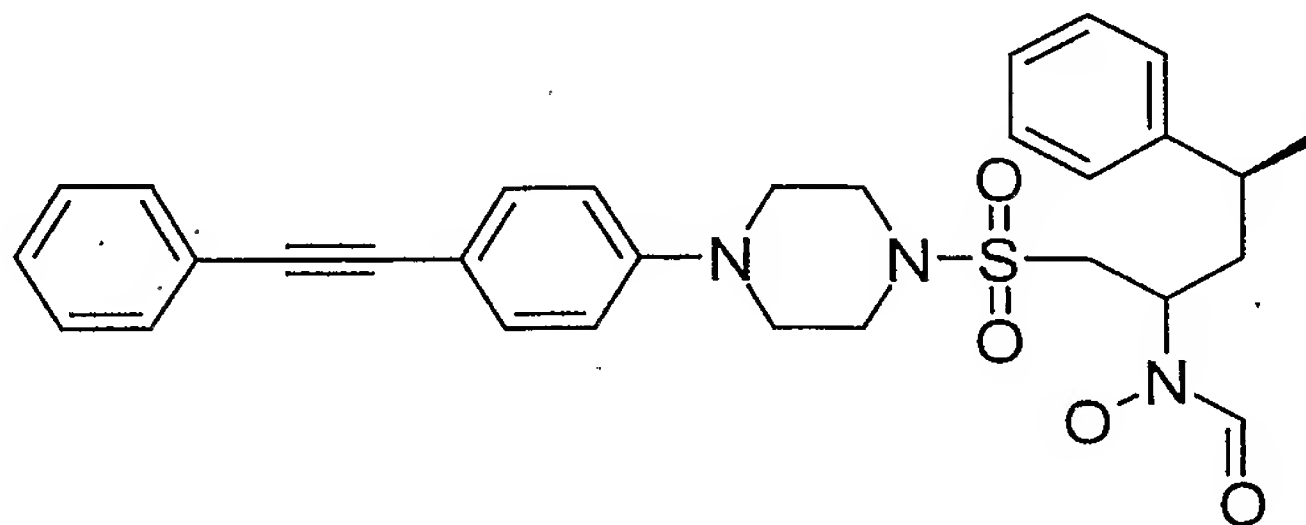
To a stirred solution of 2-((4*E/Z*)-5-{[4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine (235 mg, 0.43 mmol), in THF (5 mL) at RT was added a solution of hydroxylamine (0.5 mL, 50% aqueous solution in water). The reaction was stirred for 3 hours at RT before being  
 10 quenched with saturated aqueous ammonium chloride solution (5 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 10 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica, 10% MeOH in ethyl acetate) to give  
 15 2-(4-(hydroxyamino)-5-{[4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazin-1-yl]sulfonyl}pentyl)pyrimidine (126 mg, 51 mmol).

MS (ESI): 576.35 (MH<sup>+</sup>)

### EXAMPLE 3

20

hydroxy{(3*S*)-3-phenyl-1-[(4-[4-(phenylethynyl)phenyl]piperazin-1-yl)sulfonyl)methyl]butyl}formamide



25

To formic acid (2.5 mL, 65 mmol) at 0 °C was added acetic anhydride (500 μL, 5.2 mmol) and the mixture was stirred at 0 °C for 10 minutes. This was added to a solution of 1-{[(4*S*)-2-(hydroxyamino)-4-phenylpentyl]sulfonyl}-4-[4-(phenylethynyl)phenyl]piperazine

(345 mg, 0.685 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (5 mL) cooled to 0 °C. The reaction was brought to RT and stirred for one hour. Volatiles were then removed *in vacuo*. The residue was then precipitated in MeOH (10 mL) and heated to 40 °C for two hours, dissolving. The solution was then cooled to RT and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 1-2% MeOH in CH<sub>2</sub>Cl<sub>2</sub>) to give the title compound as a white foam (330 mg, 0.621 mmol, 91%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : complex mixture of 4 rotameric diastereomers.

MS (ESI): 532 (MH<sup>+</sup>)

The starting material was prepared as follows:

To a solution of 1-(4-bromophenyl)piperazine hydrochloride (5.09 g, 18.3 mmol, CAS number 68104-62-1) and triethylamine (7.67 mL), in CH<sub>2</sub>Cl<sub>2</sub> was added methanesulfonyl chloride (2.83 mL, 36.3 mmol) dropwise. The mixture was stirred for one hour at RT, then CH<sub>2</sub>Cl<sub>2</sub> (100 mL) was added. The organics were washed with water (2 x 100 mL), and evaporated *in vacuo* to give a yellow solid which crystallised from ethanol and was washed with ether to give 1-(methylsulfonyl)-4-(4-bromophenyl)piperazine as a white powder (4.74 g, 81%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 7.38 (d, 2 H), 6.91 (d, 2 H), 3.21 (m, 8 H), 2.89 (s, 3 H).

MS (ESI): 318, 320 (MH<sup>+</sup> Br isotope pattern).

To a stirred suspension of 1-(methylsulfonyl)-4-(4-bromophenyl)piperazine (5.77 g, 18.1 mmol), phenyl acetylene (3.97 mL, 36.1 mmol), Copper (I) iodide (38 mg, 0.2 mmol) and triphenylphosphine (95 mg, 0.36 mmol) in piperidine (72 mL) at 20 °C was added dichlorobis(triphenylphosphine)palladium(II) (38 mg, 54 μmol). The reaction was warmed to 105 °C and stirred for 14 hours under an inert atmosphere. Volatiles were removed *in vacuo*, and the solid residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (200 mL) and washed with aqueous hydrochloric acid (2M) to pH 2. The layers were then separated and the organic extract dried, (sat. Brine and MgSO<sub>4</sub>), filtered and evaporated to dryness *in vacuo*. The residue

was triturated in CH<sub>2</sub>Cl<sub>2</sub>, and filtered to give 1-(methylsulfonyl)-4-[4-(phenylethynyl)phenyl]piperazine as a flaky white solid (1.99 g, 5.86 mmol, 32 %).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 7.5 (dd, 2 H), 7.45 (d, 2 H), 7.33, (m, 3 H), 6.86 (d, 2 H), 3.38 (d, 8  
5 H), 2.82 (s, 3 H).

MS (ESI): 341 (MH<sup>+</sup>)

To a stirred suspension of 1-(methylsulfonyl)-4-[4-(phenylethynyl)phenyl]piperazine (511  
mg, 1.5 mmol) in THF (10 mL) at -40<sup>0</sup>C, was added dropwise a solution of LiHMDS in  
10 THF (3.3 mL, 1.0M solution, 3.3 mmol). The resulting suspension was stirred at -40<sup>0</sup>C for  
10 minutes before being treated with chlorotrimethylsilane (189 μL, 1.5 mmol). The  
solution was then maintained at -40<sup>0</sup>C for 10 minutes before being treated with a solution  
of (3*S*)-3-phenylbutanal (267 mg, 1.8 mmol, CAS-number 53531-19-4) in THF (6 mL).

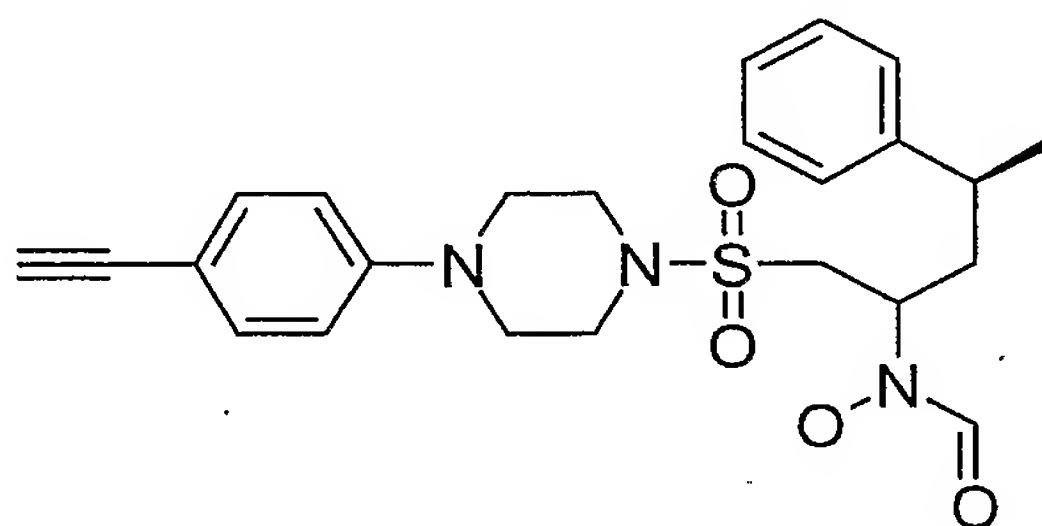
The solution was allowed to warm to -15 °C over 1 hour before being quenched with a  
15 solution of hydroxylamine (3 mL, 50% aqueous solution in water). The reaction was  
stirred for 3 hours at RT. The reaction was partitioned between ethyl acetate (30 mL) and  
water (30 mL). The combined organic extracts were then dried (sat. brine and MgSO<sub>4</sub>),  
filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography  
(silica, 35 - 55 % ethyl acetate in *i*Hexane) to give 1-{[(4*S*)-2-(hydroxyamino)-4-  
20 phenylpentyl]sulfonyl}-4-[4-(phenylethynyl)phenyl]piperazine (351 mg, 0.697 mmol).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : diastereomeric: 7.48 (m, 4 H), 7.25 (m, 8 H + CHCl<sub>3</sub>), 6.87 (q, 2 H),  
3.35 (t, 4 H), 3.23 (m, 5.5 H), 2.92 (m, 1 H), 2.76 (m, 0.5 H), 2.02 (m, 1 H), 1.83 (m, 1 H),  
1.33 (d, 3 H).

25 MS (ESI): 504 (MH<sup>+</sup>)

**EXAMPLE 4**

**hydroxy[(3*S*)-3-phenyl-1-([4-(4-ethynylphenyl)piperazin-1-yl)sulfonyl}methyl)butyl]formamide**



To formic acid (3.75 mL, 98 mmol) at 0 °C was added acetic anhydride (750 µL, 7.8 mmol) and the mixture was stirred at 0 °C for 10 minutes. This was added to a solution of 1-{[(4*S*)-2-(hydroxyamino)-4-phenylpentyl]sulfonyl}-4-[4-(trimethylsilylethynyl)phenyl]piperazine (720 mg, 94 wt %, 1.35 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (5 mL) cooled to 0 °C. The reaction was brought quickly to RT and stirred for 20 minutes. Volatiles were then removed *in vacuo*, and the residue azeotroped with toluene (3 x 5 mL). The residue was then dissolved in MeOH (10 mL) and heated to 40 °C for two hours. The solution was then cooled to RT and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 40 - 85% EtOAc in *i*Hexane) and evaporated *in vacuo*. The solid was dissolved in ether/ CH<sub>2</sub>Cl<sub>2</sub> and the insoluble contaminant removed by filtration to give the title compound as a pink foam (90 mg, 0.171 mmol, 15%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : complex mixture of 4 rotameric diastereomers.

MS (ESI): 456 (MH<sup>+</sup>)

The starting material was prepared as follows:

To a stirred suspension of 1-(methylsulfonyl)-4-(4-bromophenyl)piperazine (4.79 g, 15 mmol, prepared as in example 3), trimethylsilylacetylene (4.15 mL, 30 mmol), Copper (I) iodide (31 mg, 0.17 mmol) and triphenylphosphine (80 mg, 0.3 mmol) in piperidine (60 mL) at 20 °C was added dichlorobis(triphenylphosphine)palladium(II) (31 mg, 45 µmol).



The reaction was warmed to 105 °C and stirred for 8 hours under an inert atmosphere. After cooling, the suspension was filtered and washed with toluene. The solid was dissolved in CH<sub>2</sub>Cl<sub>2</sub>, washed with water and filtered through silica, washing with CH<sub>2</sub>Cl<sub>2</sub>. The filtrate was concentrated and the resulting suspension filtered, washing well with  
5 CH<sub>2</sub>Cl<sub>2</sub>/*i*Hexane to give 1-(methylsulfonyl)-4-[4-(trimethylsilylethynyl)phenyl]piperazine as a white powder (3.05 g, 9.06 mmol, 60 %).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 7.3 (d, 2 H), 6.9 (d, 2 H), 3.33 (t, 4 H), 3.22 (t, 4 H), 2.9 (s, 3 H), 0.2 (s, 9H).

10 MS (ESI): 337 (MH<sup>+</sup>)

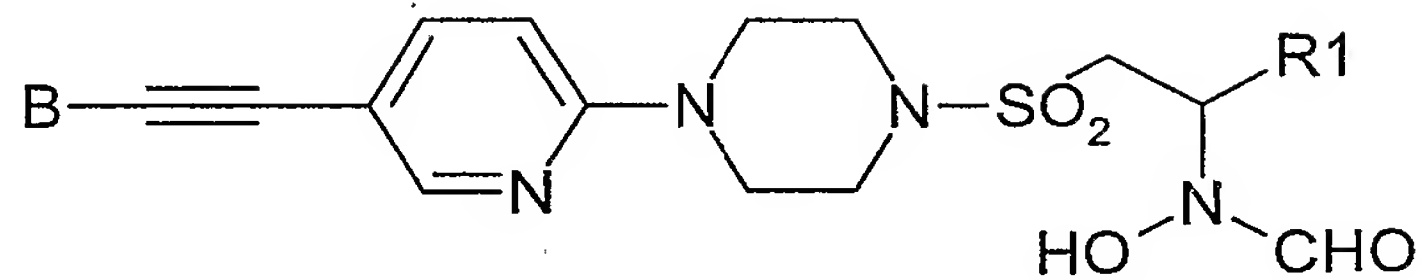
To a stirred suspension of 1-(methylsulfonyl)-4-[4-(trimethylsilylethynyl)phenyl]piperazine (673 mg, 2 mmol) in THF (13 mL) at -40<sup>0</sup>C, was added dropwise a solution of LiHMDS in THF (4.5 mL, 1.0M solution, 4.5 mmol). The  
15 resulting suspension was stirred at -40<sup>0</sup>C for 10 minutes before being treated with chlorotrimethylsilane (254 µL, 2 mmol). The solution was then maintained at -40<sup>0</sup>C for 10 minutes before being treated with a solution of (3*S*)-3-phenylbutanal (385 mg, 2.6 mmol, CAS number 53531-19-4) in THF (8 mL). The solution was allowed to warm to -15 °C over 1 hour before being quenched with a solution of hydroxylamine (1.23 mL, 50%  
20 aqueous solution in water, 20 mmol). The reaction was stirred for 3 hours at RT. The reaction was partitioned between ethyl acetate (30 mL) and water (30 mL). The combined organic extracts were then dried, (sat. brine and MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica, 30 - 40 % ethyl acetate in *i*Hexane) to give 1-{[(4*S*)-2-(hydroxyamino)-4-phenylpentyl]sulfonyl}-4-[4-(trimethylsilylethynyl)phenyl]piperazine (720 mg, 1.44 mmol, 68 % @ 94 wt %).  
25

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : diastereomeric: 7.39 (d, 2 H), 7.27 (m, 5 H + CHCl<sub>3</sub>), 6.82 (q, 2 H), 3.37(b, 4.5 H), 3.25 (b, 3 H), 3.18 (b, 2 H), 2.92 (m, 1 H), 2.75 (m, 0.5 H), 2.02 (m, 1 H), 1.83 (m, 1 H), 1.33 (d, 3 H), 0.26 (s, 9H).

30 MS (ESI): 500 (MH<sup>+</sup>)

**EXAMPLE 5**

The following compounds were also prepared.

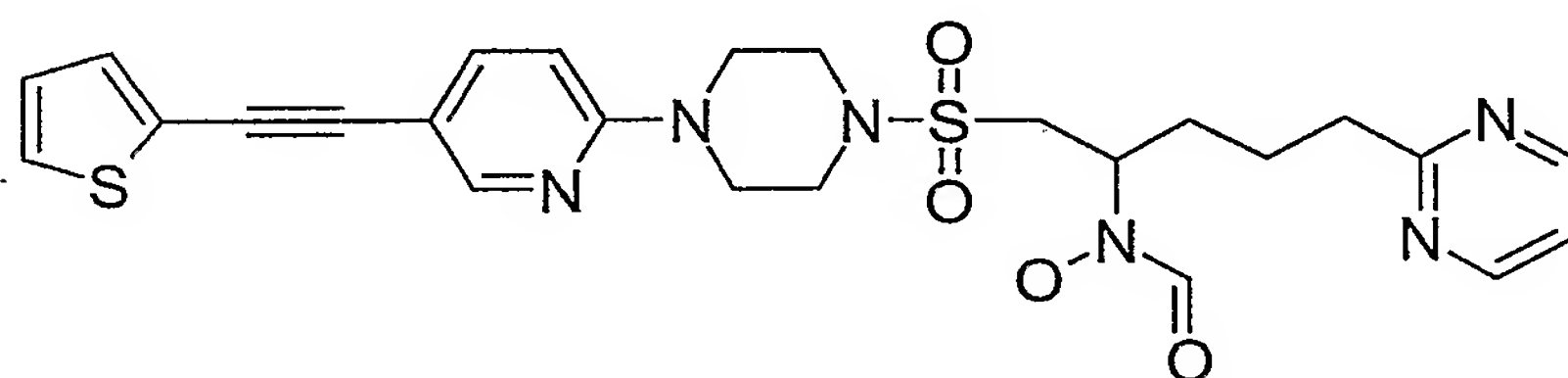


5

B	R1	M+H	Prepared using method in example
H	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	531.6	2
2-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	536.4	1
<i>tert</i> -butyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	515.5	1
<i>n</i> -propyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	501.5	1
2-Pyridyl	5-F-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	539.8	1
5-(CF <sub>3</sub> )-2-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	604.4	2
2-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	522.1	1
3-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	536.3	1
4-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	536.3	1
2-Pyrimidinyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	537.5	7
2-Pyridyl	5-Cl-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	556.21	1

**EXAMPLE 6**

**hydroxy{4-pyrimidin-2-yl-1-[(4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonyl)methyl]butyl}formamide**



To formic acid (0.3 mL, 8 mmol) at 0 °C was added acetic anhydride (75 µL, 0.8 mmol) and the mixture was stirred at RT for 10 minutes. The reaction was then recooled to 0 °C, and a solution of 2-[4-(hydroxyamino)-5-({4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pentyl]pyrimidine (84 mg, 0.16 mmol) in THF (8 mL) was then added. The reaction was brought to RT and stirred for one hour. Volatiles were then removed *in vacuo*, and the residue azeotroped with toluene (2 x 5 mL). The residue was then dissolved in MeOH (10 mL) and heated to 40 °C for one hour. The solution was then cooled to RT and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 10% MeOH in EtOAc) to give the title compound as an off-white foam (87 mg, 0.16 mmol, 54%).

<sup>1</sup>H NMR (DMSO-D<sub>6</sub>, 373K) : 9.38 (br s, 1 H), 8.68 (d, 2H), 8.30 (d, 1 H), 8.12 (br s, 1 H), 7.64 (dd, 1 H), 7.55 (d, 1 H), 7.28 (dd, 1 H), 7.22 (t, 1H), 7.09 (t, 1H), 6.85 (d, 1 H), 3.68 (m, 4 H), 3.41 (dd, 1 H), 3.23 (m, 4 H), 3.12 (dd, 1 H), 2.85 (m, 3 H), 1.70 (m, 4 H)

MS (ESI): 541.46 (MH<sup>+</sup>)

The starting material was prepared as follows:

To a stirred solution of 2-chloro-5-iodo-pyridine (CAS number 69045-79-0, 10.52 g, 43.9 mmol) and diisopropylethylamine (11.5 mL, 65.9 mmol) in DMA (200 mL) was added piperazine (15.14 g, 0.176 mol). The mixture was the heated to 120°C for 20 hours. The solid precipitate was then filtered off, and the filtrate evaporated *in vacuo*. The residue was partitioned between EtOAc (100 mL) and water (100 mL) and the layers were

separated. The aqueous layer was then extracted with EtOAc (2 x 100 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*, to give 1-(5-iodopyridin-2-yl)piperazine as a yellow solid (10.42 g, 36 mmol, 82%).

5 <sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.30 (d, 1 H), 7.65 (dd, 1 H), 6.45 (d, 1 H), 3.46, (t, 4 H), 2.97 (t, 4 H).  
MS (ESI): 290.29 (MH<sup>+</sup>)

To a stirred solution of 1-(5-iodopyridin-2-yl)piperazine (CAS number 219635-89-9, 5.083 g, 17.6 mmol), and triethylamine (4.91 mL, 35.2 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (75 mL) at 0°C  
10 was added methane sulfonyl chloride (2.04 mL, 26.4 mmol) dropwise and the resulting suspension stirred for 30 mins at room temperature. The reaction mixture was diluted with CH<sub>2</sub>Cl<sub>2</sub> (200 mL) and washed twice with water (2 x 200 mL). The organic phase was washed with saturated brine solution (1 x 200 mL), dried (MgSO<sub>4</sub>), filtered and concentrated *in vacuo* to give a dark yellow solid. This was triturated with ether, filtered  
15 and dried to give 1-(5-iodopyridin-2-yl)-4-(methanesulfonyl)piperazine as a yellow solid (5.378 g, 83%).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.33 (d, 1 H), 7.70 (dd, 1 H), 6.48 (d, 1 H), 3.63, (t, 4 H), 3.30 (t, 4 H),  
20 2.80 (s, 3 H).  
MS (ESI): 367.99 (MH<sup>+</sup>)

To a stirred suspension of 1-(5-iodopyridin-2-yl)-4-(methanesulfonyl)piperazine (7.760  
25 g, 21.1 mmol) in THF (150 mL) at -20°C, was added dropwise a solution of LiHMDS in THF (46.4 mL, 1.0M solution, 46.4 mmol). The resulting suspension was stirred at -20°C for 30 minutes before being treated with diethyl chlorophosphate (3.35 mL, 23.2 mmol). The solution was then maintained at -20°C for 15 minutes before being treated with a solution of 4-pyrimidin-2-ylbutanal (3.48 g, 23.2 mmol, CAS number 260441-10-9) in  
30 THF (20 mL). The solution was stirred at -20°C for a further 20 minutes before being quenched with saturated aqueous ammonium chloride solution (100 mL). The layers were

separated and the aqueous phase extracted with ethyl acetate (3 x 500 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. This was purified by bond-elut chromatography eluting with 25% EtOAc/hex to 50% EtOAc/hex to 100% EtOAc to give 2-((4*E/Z*)-5-{[4-(5-iodopyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine.

MS (ESI): 500.20(MH<sup>+</sup>)

To a stirred solution of 2-((4*E/Z*)-5-{[4-(5-iodopyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine (1.906 g, 3.82 mmol) in DMA (35 mL) at RT was added trimethylsilyl acetylene, (CAS number 1066-54-2, 1.08 mL, 7.64 mmol), triethylamine (1.60 mL, 11.5 mmol), CuI (291 mg, 1.53 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (442 mg, 0.38 mmol). After stirring for 30 minutes, the DMA was removed *in vacuo*, and the residue diluted with EtOAc (100 mL). The dark solution was then washed with water (100 mL). The layers were separated and the aqueous phase extracted with EtOAc (3 x 50 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 50% EtOAc in hexanes to 100% EtOAc) to give 2-{(4*E/Z*)-5-[(4-{5-[(trimethylsilyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]pent-4-enyl}pyrimidine as a yellow foam (1.609 g, 3.43 mmol, 90%). This material was used semi-crude in the next step.

To a stirred solution of 2-{(4*E/Z*)-5-[(4-{5-[(trimethylsilyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]pent-4-enyl}pyrimidine (1.605 g, 3.42 mmol) in THF (50 mL) was added TBAF (3.42 mL, 1 M solution, 3.42 mmol) and stirred at room temperature for 30 mins. The reaction mixture was then partitioned between water (200 mL) and EtOAc (200 mL). The layers were separated and the aqueous phase extracted twice with EtOAc (2 x 100 mL). The combined organics were then washed with brine (200 mL) dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was purified by bond-elut chromatography eluting with 25% EtOAc/hex to 50% EtOAc/hex to 100% EtOAc to give



2-((4*E/Z*)-5-{[4-(5-ethynylpyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine as a pale orange solid (1.065 g, 2.68 mmol, 78%).

MS (ESI): 398.10

To a stirred solution of 2-((4*E/Z*)-5-{[4-(5-ethynylpyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine (487 mg, 1.23 mmol) in DMA (20 mL) at RT was added 2-iodothiophene, (CAS number 516-12-1, 0.17 mL, 1.48 mmol), triethylamine (0.52 mL, 3.69 mmol), CuI (94 mg, 0.49 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (142 mg, 0.13 mmol). After stirring for 1 hour, the DMA was removed *in vacuo*, and the residue diluted with EtOAc (100 mL). The dark solution was then washed with water (100 mL). The layers were separated and the aqueous phase extracted with EtOAc (3 x 50 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by flash chromatography (silica gel, 50% EtOAc in hexanes to EtOAc) to give 2-[(4*E/Z*)-5-({4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pent-4-enyl]pyrimidine as a pale brown oil (371 mg, 0.69 mmol, 56%). This material was used semi-crude in the next step.

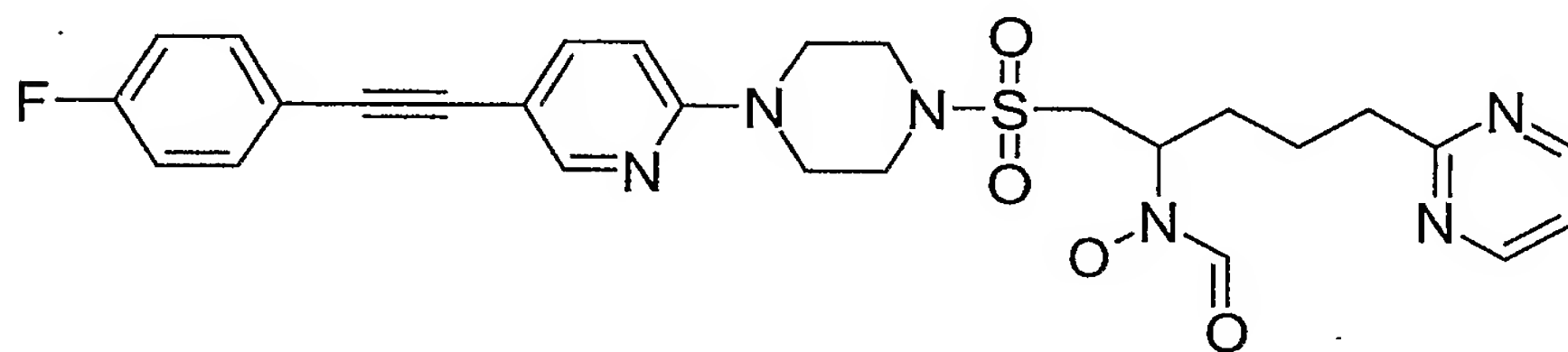
To a stirred solution of 2-[(4*E/Z*)-5-({4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pent-4-enyl]pyrimidine (371 mg, 0.69 mmol) in THF (4 mL) was added hydroxylamine (0.4 mL, 50% aqueous solution in water) and stirred for 24 hours at room temperature. The reaction was partitioned between ethyl acetate (30 mL) and water (30 mL). The combined organic extracts were then dried (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by bond elut chromatography (silica gel, EtOAc to 10% MeOH/EtOAc) to give 2-[4-(hydroxyamino)-5-({4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl}sulfonyl)pentyl]pyrimidine (84 mg, 0.16 mmol, 23%). Also recovered some isomerised SM (220mg, 0.46 mmol, 66%) presumably formed from TBAF deprotection of 2-((4*E/Z*)-5-{[4-(5-ethynylpyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine.

$^1\text{H}$  NMR ( $\text{CDCl}_3$ ) : 8.66 (m, 2 H), 8.32 (d, 1 H), 7.62 (dd, 1 H), 7.28 (m, 2 H), 7.17 (dd, 1 H), 7.02 (m, 1 H), 6.63 (m, 1 H), 5.80 (br s, 1 H), 3.74 (m, 4 H), 3.46 (m, 2 H), 3.41 (m, 4 H), 3.19 (m, 1 H), 3.04 (m, 1 H), 2.81 (m, 1 H), 2.00 (m, 2 H), 1.69 (m, 2 H).

MS (ESI): 513.41 ( $\text{MH}^+$ )

### EXAMPLE 7

1-{[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]methyl}-4-pyrimidin-2-ylbutyl(hydroxy)formamide



To formic acid (1.85 mL, 49.0 mmol) at 0 °C was added acetic anhydride (0.46 mL, 4.9 mmol) and the mixture was stirred at RT for 10 minutes. The reaction was then recooled to 0 °C, and a solution of 2-[5-[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]-4-(hydroxyamino)pentyl]pyrimidine (515 mg, 0.98 mmol) in THF (10 mL) was then added. The reaction was brought to RT and stirred for one hour. Volatiles were then removed *in vacuo*, and the residue azeotroped with toluene (2 x 5 mL). The residue was then dissolved in MeOH (10 mL) and heated to 40 °C for one hour. The solution was then cooled to RT and concentrated *in vacuo*. The white solid which precipitates upon cooling to room temperature was filtered off and dried (258 mg, 0.47 mmol, 48%).

$^1\text{H}$  NMR ( $\text{DMSO-D}_6$ , 373K) : 9.40 (br s, 1 H), 8.67 (d, 2H), 8.31 (d, 1 H), 8.12 (br s, 1 H), 7.68 (dd, 1 H), 7.55 (m, 2 H), 7.24 (t, 1 H), 7.19 (t, 2 H), 6.87 (d, 1 H), 4.34 (br s, 1 H), 3.67 (t, 4 H), 3.45 (dd, 1 H), 3.29 (t, 4 H), 3.13 (dd, 1 H), 2.89 (t, 2 H), 1.72 (m, 4 H)

MS (ESI): 553.54 ( $\text{MH}^+$ )

The starting material was prepared as follows:

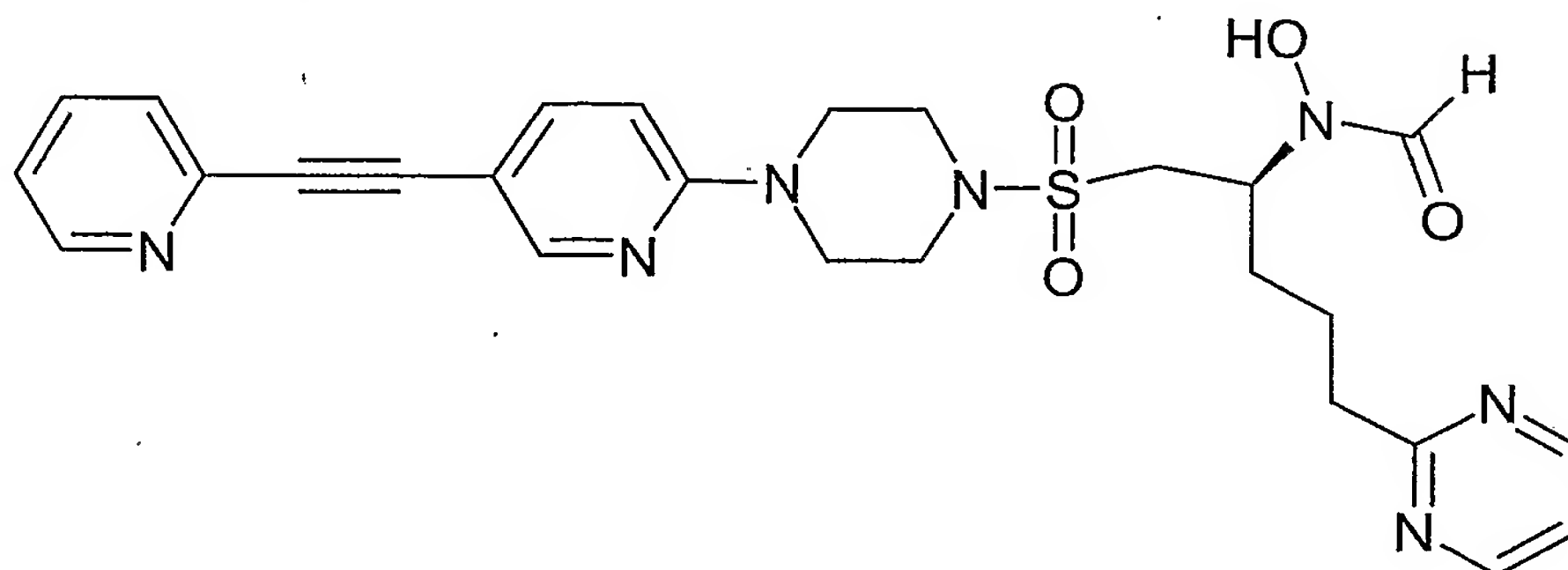
To a stirred solution of ((4*E/Z*)-5-{[4-(5-iodopyridin-2-yl)piperazin-1-yl]sulfonyl}pent-4-enyl)pyrimidine (988 mg, 1.98 mmol, prepared as in example 6) in DMA (25 mL) at RT was added 4-ethynylfluorobenzene, (CAS number 766-98-3, 476 mg, 3.96 mmol), triethylamine (0.83 mL, 5.94 mmol), CuI (151 mg, 0.8 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (229 mg, 0.2 mmol). After stirring for 60 minutes, the DMA was removed *in vacuo*, and the residue diluted with EtOAc (100 mL). The dark solution was then washed with water (100 mL). The layers were separated and the aqueous phase extracted with EtOAc (3 x 50 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by bond elut chromatography (silica gel, 50% EtOAc in hexanes to 100% EtOAc) to give 2-{(4*E/Z*)-5-[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]pent-4-enyl}pyrimidine as a pale brown solid (765 g, 1.56 mmol, 79%). This was used semi-crude in the next step.

To a solution of 2-{(4*EZ*)-5-[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]pent-4-enyl}pyrimidine (760 mg, 1.55 mmol) in THF (10 mL) was added a solution of hydroxylamine (2 mL, 50% aqueous solution in water). The reaction was stirred for 2 hours at RT before being quenched with saturated aqueous ammonium chloride solution (5 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 20 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo*. The residue was then purified by bond elut (silica, EtOAc to 10% MeOH in ethyl acetate) to give 2-[5-[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl]-4-(hydroxyamino)pentyl]pyrimidine (518 mg, 0.99 mmol, 64%).

MS (ESI): 525.33 (MH<sup>+</sup>)

**EXAMPLE 8**

hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonylmethyl]-4-pyrimidin-2-ylbutyl}formamide



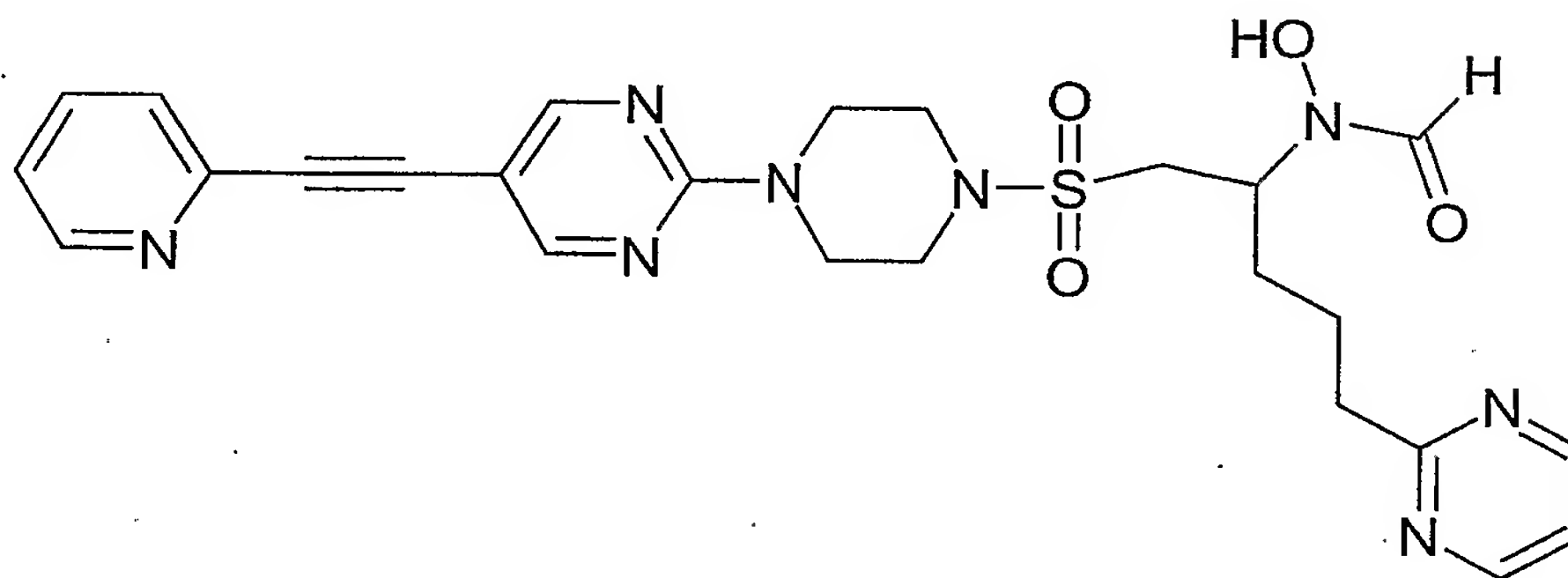
5 The racemic mixture, prepared as in example 1, was separated by chiral HPLC (on a Daicel Chiralpak AD column, 1 micron, 2cm x 25cm, 20%MeOH/MeCN eluent, flow rate 9 mL/min) to give hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonylmethyl]-4-pyrimidin-2-ylbutyl}formamide as a white solid.

10 <sup>1</sup>H NMR (DMSO-D<sub>6</sub>, 373K) : 9.41 (br s, 1 H), 8.65 (d, 2H), 8.56 (d, 1H), 8.36 (d, 1 H), 8.11 (br s, 1 H), 7.79 (m, 1 H), 7.71 (dd, 1 H), 7.55 (d, 1 H), 7.32 (m, 1 H), 7.28 (dd, 1 H), 6.88 (d, 1 H), 3.72 (m, 4 H), 3.45 (dd, 1 H), 3.32 (m, 4 H), 3.17 (dd, 1 H), 2.92 (m, 3 H), 1.77 (m, 4 H)

MS (ESI): 536.45 (MH<sup>+</sup>)

**EXAMPLE 9**

**hydroxy{1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl}formamide**



Formic acid (1.8 mL, 50 mmol) and acetic anhydride (0.5mL, 5 mmol) were mixed together at 0°C for 30 minutes, before being added to a solution 2-(4-{[2-(hydroxyamino)-5-pyrimidin-2-ylpentyl]sulfonyl}piperazin-1-yl)-5-(pyridin-2-ylethynyl)pyrimidine (190 mg, 0.37 mmol) in tetrahydrofuran (5 mL) and formic acid (1.8 mL) at 0°C. The reaction was allowed to reach room temperature and was stirred for 45 minutes. The reaction was then evaporated *in vacuo*, and azeotroped with toluene (2 x 5 mL). The residue was then dissolved in MeOH and heated to 45°C for one hour. The solution was evaporated *in vacuo*, and the residue triturated with Et<sub>2</sub>O to give a white solid which was collected by filtration, washed with Et<sub>2</sub>O and dried *in vacuo* to give hydroxy{1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl}formamide as a white solid. (64 mg, 32%).

<sup>1</sup>H NMR (d6-DMSO@373K) δ 9.50 (br, s, 1 H), 8.64 (m, 2 H), 8.61 (m, 3H), 8.17 (br, s, 1 H), 7.84 (ddd, 1 H), 7.58 (d, 1 H), 7.38 (m, 1 H), 7.29 (dd, 1 H), 4.33 (br, s, 1 H), 3.96 (m, 4 H), 3.48 (dd, 1 H), 3.33 (m, 4 H), 3.20 (dd, 1 H), 2.96 (m, 2 H), 1.74 (m, 4H).  
MS (ESI): 537.07 (MH<sup>+</sup>)

The starting material was prepared as follows :



To a stirred solution of 5-iodo-2-piperazin-1-ylpyrimidine (7.0 g, 24.1 mmol, CAS number 95847-41-9) and triethylamine (10 mL, 72 mmol) in dichloromethane (70 mL) at 0°C was added methanesulfonyl chloride (1.93 mL, 138 mmol) dropwise over 10 minutes. The reaction was then stirred for 30 minutes at 0°C, before being allowed to warm to room temperature and stirred for an additional 30 minutes. The reaction was then quenched with water (70 mL) and the layers were separated. The organic phase was washed with water (100 mL) and the organics were dried (MgSO<sub>4</sub>), filtered and evaporated *in vacuo*. The residue was then triturated with ethyl acetate and the solid residue filtered and dried *in vacuo* to give 5-iodo-2-[4-(methanesulfonyl)piperazin-1-yl]pyrimidine as an off white solid (8.85g, 24 mmol, 99%).

<sup>1</sup>H NMR (d6-DMSO) : 8.54 (s, 2 H), 3.82 (m, 4 H), 3.17, (m, 4 H), 2.87 (s, 3 H)  
MS (ESI): 369.01 (MH<sup>+</sup>)

To a stirred suspension of 5-iodo-2-[4-(methanesulfonyl)piperazin-1-yl]pyrimidine (5.52g, 15 mmol) in THF (50 mL) at -10°C, was added dropwise a solution of LiHMDS in THF (31.5 mL, 1.0M solution, 31.5 mmol). The resulting suspension was stirred at -10°C for 20 minutes before being treated with diethyl chlorophosphate (2.24 mL, 15.5 mmol). The solution was then maintained at -20°C for 20 minutes before being treated with a solution of 4-pyrimidin-2-ylbutanal (2.31 g, 15.5 mmol) in THF (5 mL). The solution was then maintained at -20 °C for one hour before being quenched with saturated aqueous ammonium chloride solution (50 mL). The layers were separated and the aqueous phase extracted with ethyl acetate (3 x 30 mL). The combined organic extracts were then dried, (MgSO<sub>4</sub>), filtered and concentrated *in vacuo* to give a brown solid which was purified by flash chromatography (silica gel, 20% to 50% to 100% EtOAc in hexanes) to give 5-iodo-2-(4-[(1*E/Z*)-5-pyrimidin-2-ylpent-1-enyl]sulfonyl)piperazin-1-ylpyrimidine as a yellow solid (5.14g, 68%, E:Z 1.7:1).

<sup>1</sup>H NMR (CDCl<sub>3</sub>) : 8.58 (m, 2 H), 8.33 (m, 2 H), 7.18, (m, 1 H), 6.73 (ddd, 1 H), 6.42 (ddd)\*, 6.04 (d, 1H), 5.93 (d)\*, 3.82 (m, 4H), 3.11 (m, 4H), 2.94 (m, 2H), 2.63 (m)\*, 2.29 (m, 2H), 1.94 (m, 2H)

\* minor geometrical isomer.

MS (ESI): 501.26 (MH<sup>+</sup>).

A stirred solution of 5-iodo-2-(4-{[(1*E*/*Z*)-5-pyrimidin-2-ylpent-1-enyl]sulfonyl}piperazin-  
5 1-yl)pyrimidine, (500 mg, 1 mmol), Pd(PPh<sub>3</sub>)<sub>2</sub>Cl<sub>2</sub> (35 mg, 0.05 mmol), CuI (76 mg, 0.4  
mmol), triethylamine (0.4 mL, 3 mmol) and 2-ethynylpyridine (206 mg, 2 mmol) in THF  
(20 mL) was stirred at room temperature for 3 hours. The mixture was then evaporated *in*  
*vacuo*, and the residue purified by flash chromatography (silica gel, 1% to 2% MeOH in  
EtOAc) to give 5-(pyridin-2-ylethynyl)-2-(4-{[(1*E*/*Z*)-5-pyrimidin-2-ylpent-1-  
10 enyl]sulfonyl}piperazin-1-yl)pyrimidine as a white solid (224 mg, 47%).

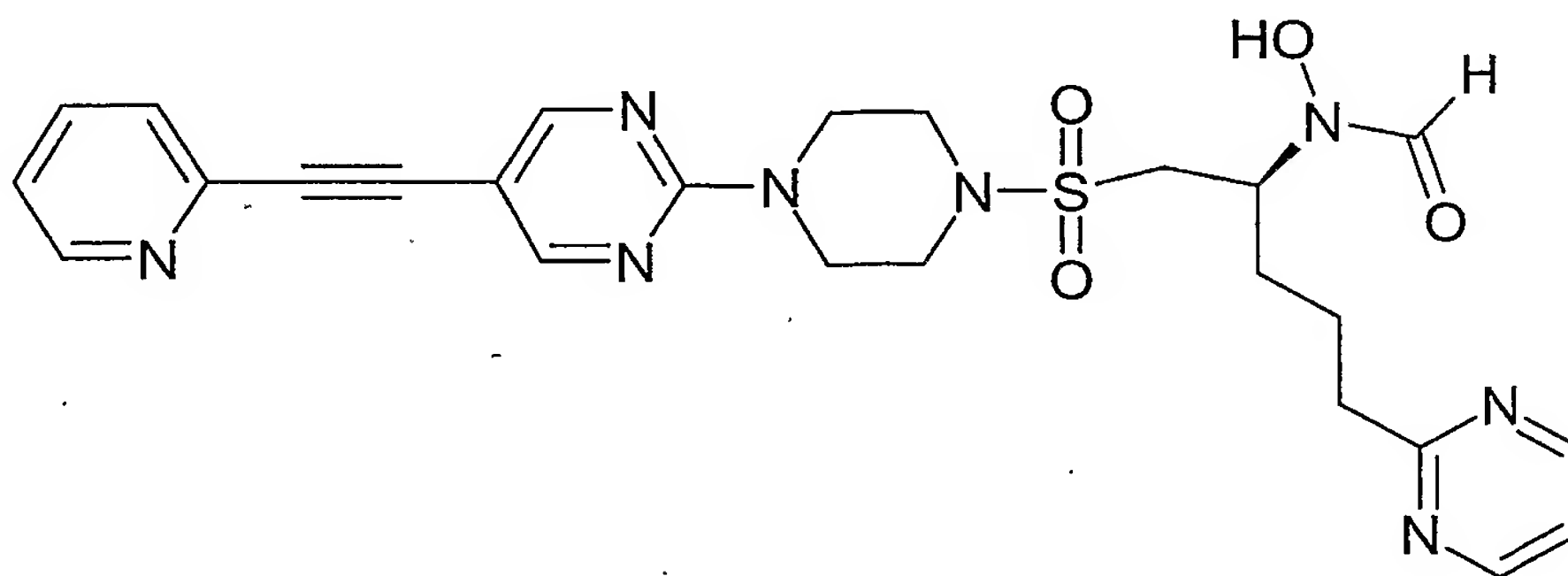
MS (ESI): 476.39 (MH<sup>+</sup>)

To a stirred solution of give 5-(pyridin-2-ylethynyl)-2-(4-{[(1*E*/*Z*)-5-pyrimidin-2-ylpent-1-  
15 enyl]sulfonyl}piperazin-1-yl)pyrimidine (220 mg, 0.46 mol) in THF (10 mL) at room  
temperature was added 50% aqueous hydroxylamine (2 mL) and the mixture stirred rapidly  
for 2 hours. The reaction was quenched by the addition of saturated ammonium chloride  
solution (5 mL) and the layers were then separated. The aqueous phase was then extracted  
with EtOAc (3 x 5 mL) and the combined organic extracts were then dried (MgSO<sub>4</sub>),  
20 filtered and evaporated *in vacuo*. The white solid obtained was then purified by flash  
chromatography (silica gel, 50% to 100% EtOAc in hexanes), to give 5 2-(4-{[2-  
(hydroxyamino)-5-pyrimidin-2-ylpentyl]sulfonyl}piperazin-1-yl)-5-(pyridin-2-  
ylethynyl)pyrimidine as a white solid (195 mg, 0.385 mmol, 83%).

25 MS (ESI): 507.06 (MH<sup>+</sup>)

**EXAMPLE 10**

**hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl}formamide**



5

The racemic mixture, prepared as in example 9, was separated by chiral HPLC (on a Merck Chiralpak AS-V column, 20  $\mu$ m, 5cm x 25cm, flow rate 50 mL / min, eluent = 100% MeOH) to give hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl}formamide as a white solid.

10

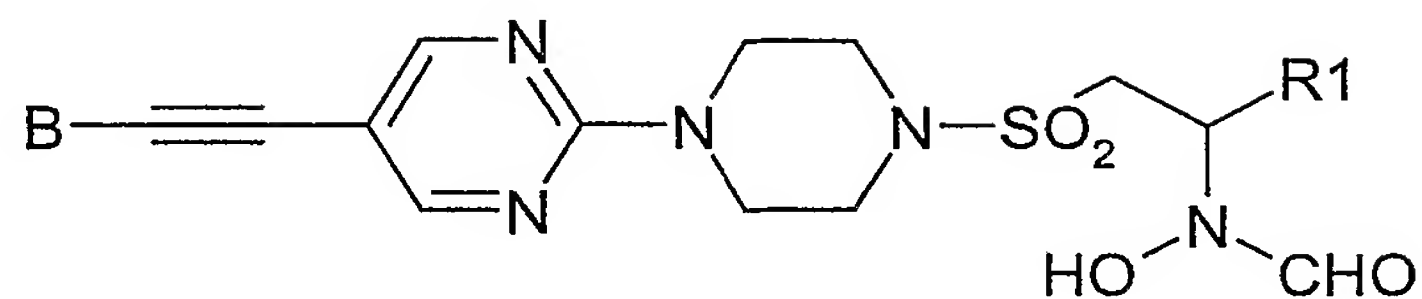
$^1\text{H}$  NMR (d6-DMSO@373k)  $\delta$  9.50 (br, s, 1 H), 8.64 (m, 2 H), 8.61 (m, 3H), 8.17 (br, s, 1 H), 7.84 (ddd, 1 H), 7.58 (d, 1 H), 7.38 (m, 1 H), 7.29 (dd, 1 H), 4.33 (br, s, 1 H), 3.96 (m, 4 H), 3.48 (dd, 1 H), 3.33 (m, 4 H), 3.20 (dd, 1 H), 2.96 (m, 2 H), 1.74 (m, 4H).

MS (ESI): 537.07 ( $\text{MH}^+$ )

15

**EXAMPLE 11**

The following compounds were also prepared.

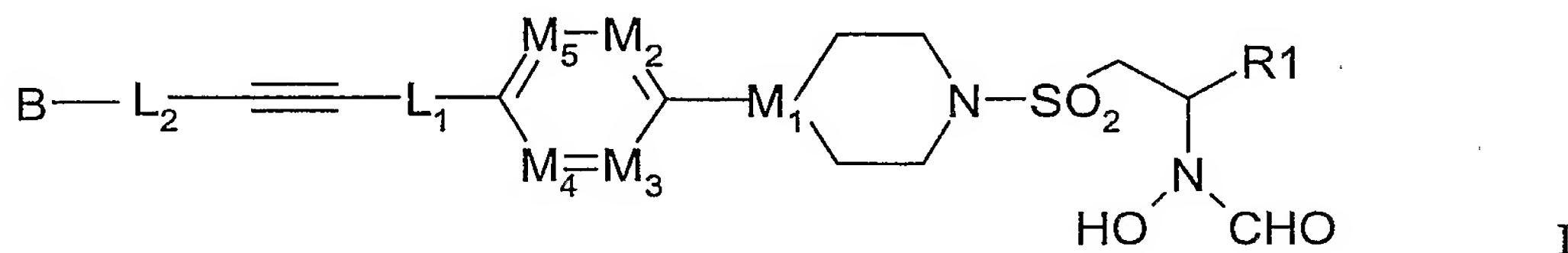


B	R1	M+H	Prepared using method in example
4-F-Ph	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	554.40	9
4-Cl-Ph	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	570.39	9
3-Pyridyl	2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub>	537.39	9
3-Pyridyl	5-F-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	541.36	9
4-F-Ph	5-F-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	558.32	9
2-Pyridyl	5-F-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	541.37	9
4-Cl-Ph	5-F-2-PyrimidinylCH <sub>2</sub> CH <sub>2</sub>	574.20	9

**CLAIMS:**

What we claim is:

1. A compound of the formula I or a pharmaceutically acceptable salt or an in vivo  
hydrolysable ester thereof



wherein

**B** is selected from H, C<sub>1-6</sub>alkyl, up to C12 cycloalkyl, up to C12 aryl, and up to C12 heteroaryl;

**B** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy;

**L<sub>1</sub>** and **L<sub>2</sub>** are each independently selected from a direct bond and C<sub>1-6</sub>alkyl;

**M<sub>1</sub>**, **M<sub>2</sub>**, **M<sub>3</sub>**, **M<sub>4</sub>** and **M<sub>5</sub>** are each independently selected from N and C;

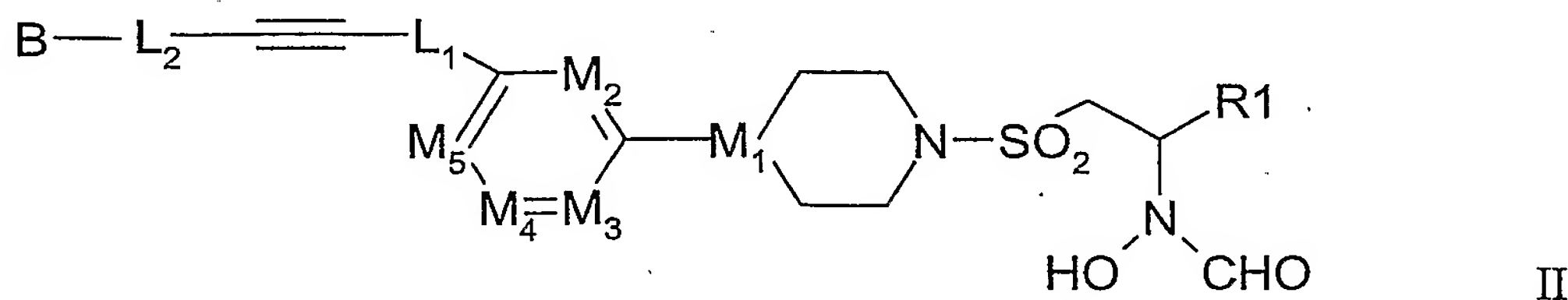
**R<sub>1</sub>** is the group -X-Y;

**X** is C<sub>1-6</sub>alkyl;

**Y** is selected from up to C10 cycloalkyl, up to C10 aryl, and up to C10 heteroaryl;

**Y** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy.

2. A compound of the formula II or a pharmaceutically acceptable salt or an in vivo  
hydrolysable ester thereof



wherein

**B** is selected from H, C<sub>1-6</sub>alkyl, up to C12 cycloalkyl, up to C12 aryl, and up to C12 heteroaryl;



**B** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy;

**L**<sub>1</sub> and **L**<sub>2</sub> are each independently selected from a direct bond and C<sub>1-6</sub>alkyl;

**M**<sub>1</sub>, **M**<sub>2</sub>, **M**<sub>3</sub>, **M**<sub>4</sub> and **M**<sub>5</sub> are each independently selected from N and C;

**R**<sub>1</sub> is the group -X-Y;

**X** is C<sub>1-6</sub>alkyl;

**Y** is selected from up to C10 cycloalkyl, up to C10 aryl, and up to C10 heteroaryl;

**Y** is optionally substituted by up to three groups independently selected from OH, NO<sub>2</sub>, CF<sub>3</sub>, CN, halogen, SC<sub>1-4</sub>alkyl, SOC<sub>1-4</sub>alkyl, SO<sub>2</sub>C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkyl, C<sub>1-4</sub>alkoxy.

3. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein **B** is selected from H, C<sub>1-6</sub>alkyl, C<sub>6</sub> aryl, and up to C<sub>6</sub> heteroaryl.

4. A compound as claimed in claim 3 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein **B** is selected from H, C<sub>2-4</sub>alkyl, C<sub>6</sub> aryl, and up to C<sub>6</sub> heteroaryl.

5. A compound as claimed in claim 4 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein **B** is up to C<sub>6</sub> heteroaryl.

6. A compound as claimed in claim 1 or claim 2 or claim 3 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein **B** is not substituted or **B** is substituted by at least one group selected from CF<sub>3</sub>, CN, halogen, C<sub>1-4</sub>alkyl.

7. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein at least one of **L**<sub>1</sub> and **L**<sub>2</sub> is a direct bond.

8. A compound as claimed in claim 7 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein each of **L**<sub>1</sub> and **L**<sub>2</sub> is a direct bond.

9. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $M_1$  is N.

10. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein at least one of  $M_2$ ,  $M_3$ ,  $M_4$ ,  $M_5$  is C, and at least one of  $M_2$ ,  $M_3$ ,  $M_4$ ,  $M_5$  is N.

11. A compound as claimed in claim 10 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein each of  $M_4$  and  $M_5$  is C, and at least one of  $M_2$  and  $M_3$  is N.

12. A compound as claimed in claim 11 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $M_2$  is C or N and  $M_3$  is N.

13. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $X$  is  $C_{2-5}$ alkyl.

14. A compound as claimed in claim 13 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $X$  is  $C_{2-3}$ alkyl.

15. A compound as claimed in claim 1 or claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $Y$  is  $C_6$  aryl or  $C_6$  heteroaryl.

16. A compound as claimed in claim 15 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $Y$  is selected from phenyl, pyridyl, pyrimidinyl, or pyrazinyl.

17. A compound as claimed in claim 1 or claim 2 or claim 15 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein  $Y$  is not substituted or  $Y$  is substituted by at least one halogen group.

18. A compound as claimed in claim 1 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein the compound of the formula I is as exemplified herein.

19. A compound as claimed in claim 18 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof wherein the compound is selected from hydroxy{-1-[(4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl} formamide, hydroxy[4-pyrimidin-2-yl-1-({4-(5-{[5-(trifluoromethyl)pyridin-2-yl]ethynyl}pyridin-2-yl)piperazin-1-yl)sulfonyl)methyl]butyl]formamide, hydroxy{(3*S*)-3-phenyl-1-[(4-[4-(phenylethynyl)phenyl]piperazin-1-yl)sulfonyl)methyl]butyl} formamide, hydroxy[(3*S*)-3-phenyl-1-({4-(4-ethynylphenyl)piperazin-1-yl)sulfonyl)methyl]butyl]formamide, hydroxy{4-pyrimidin-2-yl-1-[(4-[5-(thien-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonyl)methyl]butyl} formamid, 1-{[(4-{5-[(4-fluorophenyl)ethynyl]pyridin-2-yl}piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl(hydroxy)formamide, hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyridin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl} formamide, hydroxy{1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl} formamide, and hydroxy{(1*S*)-1-[(4-[5-(pyridin-2-ylethynyl)pyrimidin-2-yl]piperazin-1-yl)sulfonyl)methyl]-4-pyrimidin-2-ylbutyl} formamide.

20. A pharmaceutical composition which comprises a compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof and a pharmaceutically acceptable carrier.

21. A compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or in vivo hydrolysable ester thereof for use in a method of therapeutic treatment of the human or animal body.

22. A compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or in vivo hydrolysable ester thereof for use as a therapeutic agent.

5 23. A method of treating a metalloproteinase mediated disease or condition which comprises administering to a warm-blooded animal a therapeutically effective amount of a compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or in vivo hydrolysable ester thereof.

10 24. A method of treating a metalloproteinase mediated disease or condition as claimed in claim 23 which comprises treating a disease or condition mediated by MMP13.

15 25. The use of a compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or in vivo hydrolysable precursor thereof in the preparation of a medicament for use in the treatment of a disease or condition mediated by one or more metalloproteinase enzymes.

20 26. The use of a compound of the formula I as claimed in claim 1 or a compound of the formula II as claimed in claim 2 or a pharmaceutically acceptable salt or in vivo hydrolysable precursor thereof in the preparation of a medicament for use in the treatment of arthritis.

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01436

## A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 401/12, C07D 401/14, C07D 295/26, A61K 31/496, A61K 31/506, A61P 19/02  
According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CHEM. ABS. DATA

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 0012478 A1 (ZENECA LIMITED), 9 March 2000 (09.03.00) --	1-26
X	US 6100266 A (MONTANA ET AL), 8 August 2000 (08.08.00) --	1-26
P,X	WO 0162742 A1 (ASTRAZENECA AB), 30 August 2001 (30.08.01) --	1-26
P,X	WO 0162751 A1 (ASTRAZENECA AB), 30 August 2001 (30.08.01) --	1-26

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

\* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance  
 "E" earlier application or patent but published on or after the international filing date  
 "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)  
 "O" document referring to an oral disclosure, use, exhibition or other means  
 "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

25 November 2002

Date of mailing of the international search report

26-11-2002

Name and mailing address of the ISA/  
 Swedish Patent Office  
 Box 5055, S-102 42 STOCKHOLM  
 Facsimile No. +46 8 666 02 86

Authorized officer

SOLVEIG GUSTAVSSON/BS  
 Telephone No. +46 8 782 25 00



## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01436

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 0187870 A1 (DARWIN DISCOVERY LIMITED), 22 November 2001 (22.11.01)  -- -----	1-26

# INTERNATIONAL SEARCH REPORT

International application No.  
**PCT/SE02/01436**

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: **23-24**  
because they relate to subject matter not required to be searched by this Authority, namely:  
**see next sheet**
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.

Claims 23-24 relate to methods of treatment of the human or animal body by surgery or by therapy/diagnostic methods practised on the human or animal body/Rule. 39.1.(iv)). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds/compositions.

## INTERNATIONAL SEARCH REPORT

Information on patent family members

28/10/02

International application No.

PCT/SE 02/01436

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